Access DB# 107134

SEARCH REQUEST FORM

Scientific and Technical Information Center

Requester's Full Name: 404 Art Unit: 1745 Phone I	NAPLES	Examiner # :	62294	Date:	30/03
Art Unit: 1745 Phone Mail Box and Bldg/Room Location	Number 30 <u>8-1795</u> n: <i>CPU47-BE12</i> Resu	Serial No lits Format Pre	umber: <u>09/</u> ferred (circle):	432 334 PAPER DIS	K E-MAIL
If more than one search is submitted, please prioritize searches in order of need.					
Please provide a detailed statement of the Include the elected species or structures, butility of the invention. Define any terms known. Please attach a copy of the cover	search topic, and describe a reywords, synonyms, acron that may have a special me	as specifically as payms, and registry caning. Give exar	possible the subje numbers, and co	ect matter to be mbine with the	searched. concept or
Title of Invention: ENTA 64	STORAGE & GONVE	KSION DEVI	LES LISING	THERMAR	SPRANGO
Inventors (please provide full names):	RONALD GL	IIDOTTI;	Hui YE,	PENTA	
	TONGSAN	XIAD G	AL		
Earliest Priority Filing Date:	11/2/1998				
For Sequence Searches Only Please inclu appropriate serial number.	de all pertinent information (parent, child, divisi	ional, or issued pat	ent numbers) ald	ong with the
	-				
	e.				
					•
# 10 A.T.	E MATERIA =	Fos. Co	C A/5	Mi C. a	- Mos
Exs. ACTIV	E MATERIA =	1632, 60-	2, 00 32,	/013 ₂ ~	7.0001
e.					
e de la companya de La companya de la co		,			
and the second second					
State of the first state of the				<i>3</i>	
****	*******	******	*****	*****	****
STAFF USE ONLY	Type of Search	Vendo	ors and cost whe	re applicable	
Searcher:	NA Sequence (#)	STN \$	184.63		***************************************
Searcher Phone #:	AA Sequence (#)	Dialog			
Searcher Location:	Structure (#)	3 Duestel/Orbit			
Date Searcher Picked Up:	Bibliographic (a	MDKljink			
Date Completed: 10-30-03	Litigation	Lexis/Nexis			·
Searcher Prep & Review Time:	Fulltext	Sequence Systems			
Clerical Prep Time	Patent Family	WWW/Internet			
Online Time:	Other	Other (specify)			

PTO-1590 (8-01)

Access DB# 107134

SEARCH REQUEST FORM

Scientific and Technical Information Center

Requester's Full Name: JOHN MANUES Examiner #: 62294 Date: 10/36/03
Art Unit: 1745 Phone Number 30 8-1795 Serial Number: 09/432334
Requester's Full Name:
If more than one search is submitted, please prioritize searches in order of need. ***********************************
Please provide a detailed statement of the search topic, and describe as specifically as possible the subject matter to be searched. Include the elected species or structures, keywords, synonyms, acronyms, and registry numbers, and combine with the concept or utility of the invention. Define any terms that may have a special meaning. Give examples or relevant citations, authors, etc, if known. Please attach a copy of the cover sheet, pertinent claims, and abstract.
Title of Invention: ENERGY STORAGE + CONVERSION DEVICES USING THERMS SPAGNED
Title of Invention: ENERGY STORAGE + CONVERSION DEVICES USING THE RIME SPRANGE ELEGROPE Inventors (please provide full names): RONAIN GUIDOTTI; HUI YE, PENGE ELEGROPE
TONGSAN VIAN FT AT
Earliest Priority Filing Date: 1/2/1998
For Sequence Searches Only Please include all pertinent information (parent, child, divisional, or issued patent numbers) along with the appropriate serial number.
A electrode for an energy storage and conversion
device, comprising
a substrate; and
a layer of an active material comprising a metal sulfide, metal selenide, or
metal telluride, and having a thickness in the range from about 5 to about 114 microns
1 to don the substrate wherein the lover comprises orgater than 95% of the active

Exs. ACTIVE MATERIA = FeS2, Cos2, WS2, NIS2 or MOS2

material.

=> file reg
FILE 'REGISTRY' ENTERED AT 17:15:05 ON 30 OCT 2003
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
COPYRIGHT (C) 2003 American Chemical Society (ACS)

=> display history full l1-

```
FILE 'REGISTRY' ENTERED AT 16:36:47 ON 30 OCT 2003
                E IRON DISULFIDE/CN
L1
              1 SEA "IRON DISULFIDE"/CN
                E COBALT DISULFIDE/CN
L2
              1 SEA "COBALT DISULFIDE"/CN
                E TUNGSTEN DISULFIDE/CN
L3
              1 SEA "TUNGSTEN DISULFIDE"/CN
                E NICKEL DISULFIDE/CN
              1 SEA "NICKEL DISULFIDE"/CN
L4
                E MOLYBDENUM DISULFIDE/CN
L5
              1 SEA "MOLYBDENUM DISULFIDE"/CN
L6
           3032 SEA (M(L)S)/ELS (L) 2/ELC.SUB
L7
           2034 SEA (M(L)SE)/ELS (L) 2/ELC.SUB
L8
           2237 SEA (M(L)TE)/ELS (L) 2/ELC.SUB
     FILE 'HCA' ENTERED AT 16:41:36 ON 30 OCT 2003
L9
           1925 SEA L1
L10
            358 SEA L2
L11
           1747 SEA L3
L12
            345 SEA L4
           9913 SEA L5
L13
L14
         133738 SEA L6
L15
          41505 SEA L7
L16
          38851 SEA L8
                QUE ELECTROD## OR ANOD## OR CATHOD##
L17
                E COATINGS/CV
L18
           7701 SEA COATINGS/CV
                E COATING MATERIALS/CV
L19
         238921 SEA "COATING MATERIALS"/CV
                E COATING PROCESS/CV
L20
         106832 SEA "COATING PROCESS"/CV
L21
         240069 SEA ENERG? (2A) (STORAG? OR STORE# OR STORING# OR CONVERSIO
                N? OR CONVERT?) OR FUELCELL? OR FUEL? (2A) (CELL OR CELLS)
                OR BATTERY OR BATTERIES OR (ELECTROLY? OR ELECTROCHEM?
                OR GALVANI? OR PRIMARY OR SECONDARY OR WET OR DRY) (2A) (CE
                LL OR CELLS) OR DRYCELL? OR WETCELL?
L22
             14 SEA L9 AND L17 AND (L18 OR L19 OR L20) AND L21
              5 SEA L10 AND L17 AND (L18 OR L19 OR L20) AND L21
L23
              4 SEA L11 AND L17 AND (L18 OR L19 OR L20) AND L21
L24
L25
              2 SEA L12 AND L17 AND (L18 OR L19 OR L20) AND L21
            47 SEA L14 AND L17 AND (L18 OR L19 OR L20) AND L21
L26
L27
             0 SEA L15 AND L17 AND (L18 OR L19 OR L20) AND L21
L28
           518 SEA L15 AND L17 AND L21
```

```
FILE 'LCA' ENTERED AT 16:51:18 ON 30 OCT 2003
           6310 SEA FILM? OR THINFILM? OR COAT? OR TOPCOAT? OR OVERCOAT?
L29
                 OR LAYER? OR VENEER? OR CLAD? OR SHEATH? OR CASING# OR
                 ENCAS? OR ENSHEATH? OR OVERLAY? OR OVERLAID? OR ENVELOP?
     FILE 'HCA' ENTERED AT 16:54:38 ON 30 OCT 2003
L30
            222 SEA L28 AND L29
          89369 SEA L17(2A)L29
L31
L32
             78 SEA L30 AND L31
L33
           1116 SEA L15(3A)L29
L34
            530 SEA L15(3A)L17
L35
             47 SEA L32 AND (L33 OR L34)
L36
              1 SEA L32 AND L33 AND L34
L37
                 QUE (ELECTROD## OR CATHOD## OR ANOD##)/TI
L38
             22 SEA L35 AND L37
L39
              6 SEA L13 AND L17 AND (L18 OR L19 OR L20) AND L21
L40
              1 SEA L16 AND L17 AND (L18 OR L19 OR L20) AND L21
L41
           186 SEA L16 AND L17 AND L21
L42
             87 SEA L41 AND L29
L43
             18 SEA L42 AND (L33 OR L34)
L44
              3 SEA L43 AND L37
L45
              4 SEA L40 OR L44
     FILE 'REGISTRY' ENTERED AT 17:05:01 ON 30 OCT 2003
L46
           3725 SEA (L6 OR L7 OR L8) AND (T1 OR T2 OR T3)/PG
     FILE 'HCA' ENTERED AT 17:06:05 ON 30 OCT 2003
L47
          68240 SEA L46
L48
             35 SEA L47 AND L17 AND (L18 OR L19 OR L20) AND L21
L49
              0 SEA L48 AND (L33 OR L34)
L50
             26 SEA L48 AND L37
             13 SEA L50 NOT (L22 OR L23 OR L24 OR L25 OR L36 OR L38 OR
L51
                L39 OR L45 OR L43)
             37 SEA L22 OR L23 OR L24 OR L25 OR L36 OR L39 OR L43 OR L45
L52
L53
             19 SEA L38 NOT L52
             13 SEA L51 NOT (L52 OR L53)
27 SEA L52 AND (1907-1998/PRY OR 1907-1998/PY)
L54
L55
L56
            18 SEA L53 AND (1907-1998/PRY OR 1907-1998/PY)
L57
             8 SEA L54 AND (1907-1998/PRY OR 1907-1998/PY)
```

=> file hca

FILE 'HCA' ENTERED AT 17:15:27 ON 30 OCT 2003
USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
COPYRIGHT (C) 2003 AMERICAN CHEMICAL SOCIETY (ACS)

=> d 155 1-27 cbib abs hitstr hitind

L55 ANSWER 1 OF 27 HCA COPYRIGHT 2003 ACS on STN

```
139:103813 Energy storage and conversion
     devices using thin film oxide and nonoxide electrodes
     prepared by thermal spray. Guidotti, Renald A.; Ye, Hui; Xiao,
     Tongsan D.; Reisner, David E.; Doughty, Daniel H. (USA).
     Appl. Publ. US 2003138695 A1 20030724, 13 pp. (English). CODEN:
     USXXCO. APPLICATION: US 1999-432334 19991102. PRIORITY: US
     1998-PV106681 19981102.
     Thin electrodes produced by the mal spray techniques are
AΒ
     presented, wherein the thermal/spray feedstock comprises an active
     material and a protective barrier coating. In a particularly
     advantageous feature, the agtive material feedstock is a metal
     sulfide, metal selenide, or metal telluride which ordinarily decomps. at thermal spray temps. or which transforms to a material
     unsuitable for use as an electrode at thermal spray temps.
     The electrodes find particular utility in thermal
     batteries.
TΤ
     1317-33-5, Molybdenym sulfide mos2, uses 12013-10-4
     , Cobalt sulfide c\varphi's2 12035-51-7, Nickel sulfide nis2
     12068-85-8, Iron sulfide fes2 12138-09-9, Tungsten
     sulfide (WS2)
        (energy storage and conversion
        devices using thin film oxide and nonoxide electrodes
        prepd. by thermal spray)
RN
     1317-33-5 H¢A
     Molybdenum sulfide (MoS2) (8CI, 9CI) (CA INDEX NAME)
CN
S = M_0 = S
RN
     12013-10-4 HCA
     Cobalt sulfide (CoS2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
S = Co = S
     12035-51-7 HCA
RN
     Nickel sulfide (NiS2) (6CI, 8CI, 9CI) (CA INDEX NAME)
CN
s = Ni = s
     12068-85-8 HCA
RN
     Iron sulfide (FeS2) (8CI, 9CI) (CA INDEX NAME)
CN
S = Fe = S
RN
     12138-09-9 HCA
     Tungsten sulfide (WS2) (8CI, 9CI) (CA INDEX NAME)
CN
```

S = W = S

```
IC
     ICM H01M004-58
     ICS B05D005-12; B05D001-02
     429221000; 429223000; 429231500; 429218100; 427126100; 427427000
NCL
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
ST
     battery oxide film electrode thermal spray;
     energy conversion device oxide film
     electrode thermal spray
IT
     Ball milling
       Battery electrodes
       Electrodes
       Energy converters
       Energy storage systems
     Films
        (energy storage and conversion
        devices using thin film oxide and nonoxide electrodes
        prepd. by thermal spray)
IT
     Selenides
     Sulfides, uses
     Tellurides
        (energy storage and conversion
        devices using thin film oxide and nonoxide electrodes
        prepd. by thermal spray)
IT
     Coating process
        (plasma spraying, d.c. arc; energy storage
        and conversion devices using thin film oxide and
        nonoxide electrodes prepd. by thermal spray)
ΙT
     Coating process
        (thermal spraying; energy storage and
        conversion devices using thin film oxide and nonoxide
        electrodes prepd. by thermal spray)
IT
     Primary batteries
        (thermal; energy storage and
        conversion devices using thin film oxide and nonoxide
        electrodes prepd. by thermal spray)
IT
     1309-36-0, Pyrite, processes
        (energy storage and conversion
        devices using thin film oxide and nonoxide electrodes
        prepd. by thermal spray)
     1317-33-5, Molybdenum sulfide mos2, uses 12013-10-4
ΙT
     Cobalt sulfide cos2 12035-51-7, Nickel sulfide nis2
     12068-85-8, Iron sulfide fes2 12138-09-9, Tungsten
     sulfide (WS2)
        (energy storage and conversion
        devices using thin film oxide and nonoxide electrodes
        prepd. by thermal spray)
IT
     7704-34-9, Sulfur, uses 7782-49-2, Selenium, uses
                                                            13494-80-9,
     Tellurium, uses
        (energy storage and conversion
        devices using thin film oxide and nonoxide electrodes
        prepd. by thermal spray)
```

ANSWER 2 OF 27 HCA COPYRIGHT 2003 ACS on STN 134:240187 Mixed ionic electronic conductor coatings for redox electrodes. Visco, Steven J.; Chu, May-Ying (PolypPlus Battery Company, Inc., USA). (U.S. US 6210832 B1 20010403, 9 pp. (English). CODEN: USXXAM. APPLICATION: US 1998-145401 19980901. Disclosed is a redox electrode for a battery ABcell that has a coating to mitigate plugging by pptd. discharge The coating comprises a mixed ionic electronic conductor (MIEC) which is applied to the surface of a redox electrode The presence of the MIEC coating allows for rapid removal of discharge product ppts. from redox electrodes since it is capable of conducting both electrons and ions. As a result, the chem. action necessary to remove such ppts. may take place on both the electrolyte side of the ppt. and at the ppt./electrode interface. MIEC coatings in accordance with the present invention may be composed of any suitable material having ionic cond. for a metal ion in a neg. electrode with which the redox electrode is to be paired in a battery cell, and reversible redox capacity. Examples include TiS2, FeS2, and cobalt IT12068-85-8, Iron disulfide (mixed ionic electronic conductor coatings for redox electrodes) 12068-85-8 HCA RN CNIron sulfide (FeS2) (8CI, 9CI) (CA INDEX NAME) S == Fe == S IC ICM H01M004-02 NCL 429218100 CC52-2 (Electrochemical, Radiational, and Thermal Energy Technology) ST battery redox electrode mixed ionic electronic conductor coating ITBattery cathodes Coating materials Secondary batteries (mixed ionic electronic conductor coatings for redox electrodes) ITAlkali metal sulfides Polysulfides Sulfides, uses (mixed ionic electronic conductor coatings for redox electrodes) IT Alkali metal compounds Alkaline earth compounds

IT7704-34-9, Sulfur, uses 74432-42-1, Lithium polysulfide (mixed ionic electronic conductor coatings for redox electrodes) IT

redox electrodes)

7440-44-0, Carbon, uses 11104-61-3, Cobalt oxide 12039-13-3,

(polysulfides; mixed ionic electronic conductor coatings for

Titanium disulfide 12068-85-8, Iron disulfide (mixed ionic electronic conductor coatings for redox electrodes)

L55 ANSWER 3 OF 27 HCA COPYRIGHT 2003 ACS on STN
134:210599 Long cycle-life alkali metal battery with
cathode coated with a very thin protective film. Peled,
Emanuel; Golodnitsky, Diana; Strauss, Ela (Ramot University
Authority for Applied Research and Industrial Development L,
Israel). U.S. US 6203947 B1 20010320, 16 pp. (English). CODEN:
USXXAM. APPLICATION: US 1999-280646 19990329. PRIORITY: IL
1998-124007 19980408.

AB The present invention provides a cathode for use in a secondary electrochem. cell, such cathode being coated with a very thin, protective film, permeable to ions. The protective film of the cathode usually has a thickness of up to about 0.1 .mu.m and it provides protection against high voltage charging and overdischarging. The present invention further provides a secondary electrochem. cell comprising such a cathode.

RN 12068-85-8 HCA

CN Iron sulfide (FeS2) (8CI, 9CI) (CA INDEX NAME)

S=== S

IC ICM H01M004-58

NCL 429231950

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 38

ST battery cathode protective film coated

IT Alloys, uses

(alkali metal; long cycle-life alkali metal battery with cathode coated with very thin protective film)

IT Alkali metals, uses

(alloys; long cycle-life alkali metal battery with cathode coated with very thin protective film)

IT Fluoropolymers, uses Polycarbonates, uses Polyoxyalkylenes, uses

(binder; long cycle-life alkali metal battery with cathode coated with very thin protective film)

IT Polyoxyalkylenes, uses

(lithium complex; long cycle-life alkali metal battery with cathode coated with very thin protective film)

IT Battery cathodes
Coating materials
Polymer electrolytes

```
(long cycle-life alkali metal battery with
        cathode coated with very thin protective film)
IT
     Alkali metals, uses
        (long cycle-life alkali metal battery with
        cathode coated with very thin protective film)
IT
     9003-17-2, Polybutadiene
                                 9003-53-6, Polystyrene
                                                          24937-79-9, Pvdf
     25014-41-9, Polyacrylonitrile
                                      25322-68-3, Peo
        (binder; long cycle-life alkali metal battery with
     cathode coated with very thin protective film)
7429-90-5, Aluminum, uses 7440-02-0, Nickel, uses
IT
                                                            12597-68-1,
     Stainless steel, uses
        (current collector; long cycle-life alkali metal battery
        with cathode coated with very thin protective film)
     7439-89-6, Iron, uses 7439-95-4, Magnesium, uses
IT
                                                           7439-96-5,
     Manganese, uses
                       7440-42-8, Boron, uses
                                               7440-48-4, Cobalt, uses
     7440-70-2, Calcium, uses
        (dopant; long cycle-life alkali metal battery with
        cathode coated with very thin protective film)
IT
     1309-48-4, Magnesia, uses 1314-23-4, Zirconia, uses
                                                              1344-28-1,
     Alumina, uses
                     7631-86-9, Silica, uses
                                               13463-67-7, Titania, uses
        (filler; long cycle-life alkali metal battery with
        cathode coated with very thin protective film)
IT
     96-49-1, Ethylene carbonate
                                  105-58-8, Diethyl carbonate
     108-32-7, Propylene carbonate 616-38-6, Dimethyl carbonate
     623-53-0, Ethyl methyl carbonate 1309-36-0, Pyrite, uses
     1314-62-1, Vanadium pentoxide, uses 7439-93-2, Lithium, uses
     7550-35-8, Lithium bromide
                                 7570-02-7, DiVinyl carbonate
     10377-51-2, Lithium iodide
                                  10411-26-4, Butyl carbonate
     12031-65-1, Lithium nickel oxide linio2 12039-13-3, Titanium
     disulfide
               12057-17-9, Lithium manganese oxide limn2o4
     12068-85-8, Iron sulfide fes2
                                     12190-79-3, Cobalt lithium
     oxide colio2
                    14283-07-9, Lithium tetrafluoroborate
                                  24991-55-7, Polyethylene glycol
     Lithium hexafluorophosphate
                      25322-68-3D, Peo, lithium complex
     dimethyl ether
                                                          26098-78-2,
     Ethylene oxide-methylmethacrylate copolymer
                                                   90076-65-6
     329038-54-2, Vanadium oxide (V8013)
        (long cycle-life alkali metal battery with
        cathode coated with very thin protective film)
IT.
     7439-93-2D, Lithium, polyethylene oxide complex, uses
        (long cycle-life alkali metal battery with
        cathode coated with very thin protective film)
IT
     33454-82-9, Lithium triflate
        (stainless steel coated with; long cycle-life alkali metal
        battery with cathode coated with very thin
        protective film)
     7440-44-0, Carbon, uses 7782-42-5, Graphite, uses
IT
        (stainless steel coated with; long cycle-life alkali metal
        battery with cathode coated with very thin
        protective film)
```

ANSWER 4 OF 27 HCA COPYRIGHT 2003 ACS on STN

132:38125 Thermal sprayed electrodes for energy

storage or conversion devices. Ye, Hui; Strock, Christopher; Xiao, Tongsan; Strutt, Peter R.; Reisner, David E. (Us Nanocorp, Inc., USA). PCT Int. Appl. WO 9964641 A1 19991216, 19 pp. DESIGNATED STATES: W: CN, JP, US; RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE. (English). CODEN: PIXXD2. APPLICATION: WO 1999-US12899 19990609. (PRIORITY: US / 1998-88777 (19980610) AΒ Film electrodes are manufd. by coating of an active material feedstock powder (e.g., FeS2) with an additive material (e.g., S) suitable for preventing thermal decompn. of the feedstock powder during thermal spraying. The coated feedstock is sprayed onto a substrate using a plasma gun, forming a coating on the substrate, thereby providing an electrode. 1317-33-5, Molybdenum disulfide, uses 12013-10-4, ITCobalt disulfide 12068-85-8, Iron disulfide 12138-09-9, Tungsten sulfide (thermal sprayed film electrodes for energy storage or conversion devices) RN1317-33-5 HCA CNMolybdenum sulfide (MoS2) (8CI, 9CI) (CA INDEX NAME) S = Mo = S12013-10-4 HCA RNCobalt sulfide (CoS2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CN S = Co = SRN12068-85-8 HCA CN Iron sulfide (FeS2) (8CI, 9CI) (CA INDEX NAME) S = Fe = S12138-09-9 HCA RNTungsten sulfide (WS2) (8CI, 9CI) (CA INDEX NAME) CN S = W = SIC ICM C23C004-04 C23C004-12; H01M004-04 CC52-2 (Electrochemical, Radiational, and Thermal Energy Technology) electrode manuf thermal spraying STIT (porous; thermal sprayed film electrodes for energy storage or conversion devices) Film electrodes IT(thermal sprayed film electrodes for energy storage or conversion devices)

Maples 09/432,334 Carbonaceous materials (technological products) IT Metals, uses Oxides (inorganic), uses Sulfides, uses (thermal sprayed film electrodes for energy storage or conversion devices) IT Coating process (thermal spraying; thermal sprayed film electrodes for energy storage or conversion devices) IT 7704-34-9, Sulfur, uses 9005-25-8, Starch, uses (additive; thermal sprayed film electrodes for energy storage or conversion devices) 7429-90-5, Aluminum, uses IT 7440-32-6, Titanium, uses (substrate; thermal sprayed film electrodes for energy storage or conversion devices) IT1309-36-0, Pyrite, uses 1313-13-9, Manganese dioxide, uses 1313-99-1, Nickel oxide, uses 1317-33-5, Molybdenum disulfide, uses 1344-70-3, Copper oxide 7440-44-0, Carbon, uses 7440-44-0D, Carbon, fluorinated, uses 11104-61-3, Cobalt oxide 11105-02-5, Silver vanadium oxide 11115-78-9, Copper sulfide 11118-57-3, Chromium oxide 11126-12-8, Iron sulfide 12013-10-4, Cobalt disulfide 12039-13-3, Titanium disulfide 12054-48-7, Nickel hydroxide 12068-85-8, Iron disulfide 12138-09-9, Tungsten sulfide 181183-66-4, Copper silver vanadium oxide (thermal sprayed film electrodes for energy storage or conversion devices) ANSWER 5 OF 27 HCA COPYRIGHT 2003 ACS on STN L55 123:98745 Process for coating with single source precursors. Winter, Charles H.; Lewkebandara, T. Suren (Wayne State University, U.S. US 5425966 A 19950620, 7 pp. (English). CODEN: USXXAM. APPLICATION: US 1994-329841 19941027.

AΒ Reaction products of metal halides with org. chalcogenides and dichalcogenides provide single source precursors for metal dichalcogenide coatings. The single source precursors are sublimed at reduced pressure and allowed to contact a substrate maintained at an elevated temp. The resulting dichalcogenide coatings (for example TiS2) are smooth, and adherent, and may be used in numerous applications, for example, as cathodes for Li batteries.

IT12067-45-7, Titanium diselenide 12067-75-3, Titanium ditelluride

(film deposition using single source precursor)

RN12067-45-7 HCA

Titanium selenide (TiSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CN

Se=Ti=Se

RN12067-75-3 HCA

Titanium telluride (TiTe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CN

Te-Ti-Te IC ICM C23C016-00 NCL 427255100 CC 75-1 (Crystallography and Liquid Crystals) Section cross-reference(s): 52 STGroup IVB chalcogenide deposition single precursor; titanium chalcogenide film deposition single precursor Group IVB element compounds IT (halo org. chalcogenide complexes; prepn. and use as single source precursor for metal dichalcogenide film deposition) ΙT Vapor deposition processes (of metal dichalcogenide films using single source precursor) TIGroup IVB element halides (reaction with org. chalcogen compds. in prepn. of single source precursors for metal dichalcogenide films) ΙT 12039-13-3, Titanium disulfide 12067-45-7, Titanium diselenide 12067-75-3, Titanium ditelluride (**film** deposition using single source precursor) IT165327-11-7P 165327-12-8P (prepn. and use as single source precursor for metal dichalcogenide film deposition) IT 16893-00-8P 16893-01-9P 16920-83-5P 57965-49-8P (prepn. as single source precursor for metal dichalcogenide film deposition) IT 7550-45-0, Titanium tetrachloride, reactions (reaction with org. chalcogen compds. in prepn. of single source precursors for metal dichalcogenide films) ΙT 165327-13-9 (titanium sulfide film deposition using single source precursor of) ANSWER 6 OF 27 HCA COPYRIGHT 2003 ACS on STN L55 122:85495 Secondary alkali metal battery. Kozmik, Ivan D.; Tovstjuk, Kornei D.; Kovalyuk, Zahar D.; Grigortchak, Ivan I.; Krigan, Elvria G.; Bahmatyuk, Boqdan P. (I. N. Frantsevich Institute for Problems in Materials Science, Ukraine). U.S. US(8368957 A 19941129, 12 pp. Cont.-in-part of U.S. Ser. No.783,886, abandoned. (English). CODEN: USXXAM. APPLICATION: US 1992-968183 19921029. PRIORITY: US 1991-783886 19911029. AΒ The battery with improved capacity contains a Group IA element (G) anode, an electrolyte capable of ion transport of anode species, and a cathode comprising a

layered cryst. material GxMyXz, where x .ltoreq.10, M is Bi and Sb, y = 1 and 2, X is S, Se and Te, and z = 1, 2, and 3. The layered cryst. material has a sufficiently low defect d. and appropriate impurity distribution, together sufficient to permit intercalation of .gtoreq.3 mol Li within van der Waals channels/mol

of said material without significant distortion of the lattice. The change in Gibbs free energy of this material is substantially independent of the intercalated Li concn.

IT 1304-82-1P, Bismuth telluride 12068-69-8P, Bismuth selenide

(battery cathodes from lithium-

intercalatable)

RN 1304-82-1 HCA

CN Bismuth telluride (Bi2Te3) (8CI, 9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

RN 12068-69-8 HCA

CN Bismuth selenide (Bi2Se3) (6CI, 8CI, 9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
===========	+=====================================	<u> </u>
Se	3	7782-49-2
Bi	2	7440-69-9

IC ICM H01M004-58

NCL 429194000

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

ST alkali metal secondary battery; lithium intercalatable chalcogenide battery cathode

IT Batteries, secondary

(alkali metal)

IT Cathodes

(battery, from lithium-intercalatable antimony or bismuth chalcogenides)

IT 7439-93-2P, Lithium, uses

(battery cathodes from antimony or bismuth

chalcogenides intercalated with)

IT 1304-82-1P, Bismuth telluride 1345-07-9P, Bismuth sulfide

12068-69-8P, Bismuth selenide

(battery cathodes from lithium-

intercalatable)

IT 7440-36-0D, Antimony, chalcogenides

(battery cathodes from lithium-

intercalatable)

IT 1304-76-3, Bismuth oxide, uses 1317-38-0, Copper oxide (CuO), uses (battery cathodes from lithium-intercalatable chalcogenides contg.)

L55 ANSWER 7 OF 27 HCA COPYRIGHT 2003 ACS on STN

120:81559 Stacked cell array bipolar battery with thermal sprayed container and cell seal. Williams, Mark T.; Briscoe, James D.; Oweis, Salah M. (Saft America Inc., USA). U.S. US 5254415 X 19931019, 11 pp. (English). CODEN: USXXAM. APPLICATION: US 1992-865471 19920409.

AB A Li alloy/metal sulfide **battery** comprises a stacked array of cells spray coated with a ceramic coating followed by a metallic containment structural coating (e.g., from stainless steel SUS410)

to maintain the structural integrity of the array during high-temp. use. The ceramic layer preferably comprises Li2S, CaS, SrS, BaS, Li2O, BeO, MgO, CaO, SrO, Ba2O, Li3N, Be3N2, Mg3N2, Ca3N2, Sr3N2, Si3N4, BN, AlN, and their mixts. IT 12013-10-4, Cobalt sulfide (CoS2) 12035-51-7, Nickel sulfide (NiS2) 12068-85-8, Iron sulfide (FeS2) (cathode, in stacked cell array bipolar battery with thermal sprayed container and cell seal) 12013-10-4 HCA RN CNCobalt sulfide (CoS2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) $S = C_0 = S$ RN12035-51-7 HCA CN Nickel sulfide (NiS2) (6CI, 8CI, 9CI) (CA INDEX NAME) S = Ni = SRN12068-85-8 HCA Iron sulfide (FeS2) (8CI, 9CI) (CA INDEX NAME) CN S = Fe = SIC ICM H01M006-46 ICS H01M002-08; H01M010-18; H01M004-36 NCL 429153000 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CCST battery stacked ceramic coating; magnesia coating stacked battery ITCoating materials (ceramics, in stacked cell array bipolar battery with thermal sprayed container and cell seal) IT Ceramic materials and wares (coating, in stacked cell array bipolar battery with thermal sprayed container and cell seal) Batteries, primary IT (stacked, lithium alloy-metal sulfide, with ceramic coating, for high-temp. use) IT72785-69-4 (anode, in stacked cell array bipolar battery with thermal sprayed container and cell seal) 1314-62-1, Vanadium oxide (V2O5), uses IT 1317-42-6, Cobalt sulfide CoS) 12013-10-4, Cobalt sulfide (CoS2) 12031-65-1, Lithium nickel oxide (LiNiO2) 12035-51-7, Nickel sulfide 12039-13-3, Titanium sulfide (TIS2) 12068-85-8, Iron sulfide (FeS2) 12190-79-3, Cobalt lithium oxide (LiCoO2) 13463-67-7, Titania, uses 16812-54-7, Nickel sulfide (NiS) (cathode, in stacked cell array bipolar battery with thermal sprayed container and cell seal)

ΙT 1304-54-7, Beryllium nitride (Be3N2) 1304-56-9, Beryllium oxide (BeO) 1305-78-8, Calcia, uses 1309-48-4, Magnesia, uses 1314-11-0, Strontium oxide (SrO), uses 1314-96-1, Strontium 1314-96-1, Strontium sulfide (SrS) 7439-98-7, Molybdenum, uses 10043-11-5, Boron nitride (BN), uses 12013-82-0, Calcium nitride (Ca3N2) 12033-82-8, Strontium nitride (Sr3N2) 12033-89-5, Silicon nitride, 12057-24-8, Lithium oxide (Li20), uses 12057-71-5, Magnesium nitride (Mg3N2) 12136-58-2, Lithium sulfide (Li2S) 12231-50-4, Barium oxide (Ba20) 12611-79-9, SUS410 20548-54-3, 21109-95-5, Barium sulfide (BaS) Calcium sulfide (CaS) 24304-00-5, Aluminum nitride (AlN) 26134-62-3, Lithium nitride (Li3N)

(coating, in stacked cell array bipolar **battery** with thermal sprayed container and cell seal)

L55 ANSWER 8 OF 27 HCA COPYRIGHT 2003 ACS on STN

111:60991 Photochargeable battery. Takada, Kazunori; Kanbara,
Teruhisa; Tonomura, Tadashi; Kondo, Shigeo (Matsushita Electric
Industrial Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 01007480 A2

19890111 Heisei, 6 pp. (Japanese). CODEN: JKXXAF.

APPLICATION: JP 1987-162780 19870630.

AΒ The title battery has a solid electrolyte, a photoconductive electrode coated with a material which can electrochem. insert or exclude a mobile metal ion of the electrolyte, and a counterlectrode contq. that metal or ion. photoconductive **electrode** can be an n- or p-type orq. or inorg. semiconductor. When the mobile ion is Cu ion, the coating material can be Chevrel-type CuxMo6S8-y (x .ltoreq.8, y .ltoreq.0.5), the electrolyte can be RbCu4I2-zCl3+z (z 0.5-1), and the counterelectrode can be Cu or the above CuxMo6S8-y; when the mobile ion is Ag, the coating material can be Chevrel-type AgaMo6S8-b (a .ltoreq.8, b .ltoreq.0.5), the electrolyte can be RbAq4I2-zCl3+z, and the counterelectrode can be Ag of AgaMo6S6-b. Thus, a battery using a Cu2Mo6S8coated ITO electrode, a RbCu4I1.5Cl3.5 electrolyte, and a Cu2S counterelectrode had a higher capacity than a battery using a Au-coated ITO electrode.

1306-25-8, Cadmium telluride (CdTe), uses and miscellaneous
12058-18-3, Molybdenum selenide (MoSe2) 12067-46-8
, Tungsten diselenide

(electrodes from Chevrel-type compd.-coated,
photoconductive n-type, for photochargeable batteries)

RN 1306-25-8 HCA

CN Cadmium telluride (CdTe) (9CI) (CA INDEX NAME)

Cd=== Te

RN 12058-18-3 HCA

CN Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

```
Se Mo Se
RN
     12067-46-8 HCA
     Tungsten selenide (WSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
Se = W = Se
IC
     ICM H01M014-00
     ICS H01L031-04
CC
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
ST
     battery photochargeable photoconductive electrode
     coating; ITO electrode copper molybdenum sulfide
IT
     Electrodes
        (battery, semiconductor, Chevrel-type compd. -
        coated)
IT
     1303-00-0, Gallium arsenide (GaAs), uses and miscellaneous
     1306-23-6, Cadmium sulfide (CdS), uses and miscellaneous
     1306-24-7, Cadmium selenide (CdSe), uses and miscellaneous
     1306-25-8, Cadmium telluride (CdTe), uses and miscellaneous
     1317-33-5, Molybdenum disulfide, uses and miscellaneous
     Silicon, uses and miscellaneous 12018-95-0, Copper indium selenide
     (CuInSe2) 12058-18-3, Molybdenum selenide (MoSe2)
     12063-98-8, Gallium phosphide (GaP), uses and miscellaneous
     12064-03-8, Gallium antimonide (GaSb) 12067-46-8, Tungsten
     diselenide
                  22398-80-7, Indium phosphide (InP), uses and
                     37382-15-3, Aluminum gallium arsenide ((Al,Ga)As)
     121857-55-4, Indium silver sulfide (InAg0-1S2)
                                                      121857-78-1, Copper
     indium selenide (Cu0-1InSe2) 121857-79-2, Copper indium selenide
     (Cu3-1InS2)
        (electrodes from Chevrel-type compd.-coated,
        photoconductive n-type, for photochargeable batteries)
IT
     7681-65-4, Copper iodide (CuI) 7758-89-6, Copper chloride (CuCl)
     7783-90-6, Silver chloride (AgCl), uses and miscellaneous
     7783-96-2, Silver iodide (AgI)
                                     7785-23-1, Silver bromide (AgBr)
     7787-70-4, Copper bromide (CuBr) 121857-74-7, Copper sulfide
                121918-00-1, Silver sulfide (Ag1-2S)
     (Cu1-2S)
        (electrodes from Chevrel-type compd.-coated,
        photoconductive p-type, for photochargeable batteries)
     51912-50-6, Copper molybdenum sulfide (CuMo3S4)
IT
                                                      120922-23-8,
     Molybdenum silver sulfide (Mo3AgS4) 121857-76-9, Molybdenum silver
                               121857-77-0, Copper molybdenum sulfide
     sulfide (Mo6Aq0-8S7.5-8)
     (Cu0-8Mo6S7.5-8)
        (electrodes from semiconductors coated with
        Chevrel-type, photoconductive, for photochargeable
        batteries)
IT
     22205-45-4, Copper sulfide (Cu2S)
        (electrodes, for photochargeable batteries)
     73379-32-5, Copper rubidium chloride iodide (Cu4RbCl3.5I1.5)
IT
     121857-75-8, Rubidium silver chloride iodide (RbAg4Cl3.5-4I1-1.5)
        (electrolyte, for photochargeable batteries)
```

```
ANSWER 9 OF 27 HCA COPYRIGHT 2003 ACS on STN
109:233267 Low temperature chemical preparation of semiconducting
     transition metal chalcogenide films for energy
     conversion and storage, lubrication and surface
     protection. Chatzitheodorou, G.; Fiechter, S.; Kunst, M.; Luck, J.; Tributsch, H. (Hahn-Meitner-Inst., Berlin, Fed. Rep. Ger.).
     Materials Research Bulletin, 23(9), 1261-71 (English) 1988
        CODEN: MRBUAC. ISSN: 0025-5408.
AΒ
     A technique is presented for the prodn. of films of transition-metal
     disulfides, such as MoS2, WS2, FeS2, and RuS2, by the reaction of
     transition metal carbonyls (e.g., Mo(Co)6, W(CO)6, Fe(CO)5,
     Ru3(CO)12) with a S source (e.g., S, H2S) in an org. solvent (e.g.,
     C6H6, PhMe, xylene, mesitylene (1,3,5-trimethylbenzene)) at
     80-165.degree.. The quality of the materials and films was studied.
     Some applications are discussed: as photoactive materials (e.g.,
     MoS2, WS2, FeS2), as lubricating films (MoS2), as electrodes
     for Li batteries (MoS2, FeS2), and in corrosion protection
     (RuS2).
     12068-85-8, Iron disulfide
IT
        (deposition of film of, by reaction of iron carbonyl and sulfur
        and hydrogen sulfide and thiourea)
     12068-85-8 HCA
RN
CN
     Iron sulfide (FeS2) (8CI, 9CI) (CA INDEX NAME)
S = Fe = S
     1317-33-5P, Molybdenum disulfide, preparation
IT
        (deposition of film of, by reaction of molybdenum carbonyl and
        sulfur)
     1317-33-5 HCA
RN
CN
     Molybdenum sulfid
                                                       NAME)
S = Mo = S
     12138-09-9, Tungs
IT
        (deposition of
                                                       ten carbonyl and
        sulfur)
RN
     12138-09-9 HCA
CN
     Tungsten sulfide
                                                       ME)
S = W = S
CC
     47-10 (Apparatus
ΙT
     Batteries, primar
        (lithium-trans
                                                       tion,
        transition-met
IT
     Coating materials
        (anticorrosive, ruthenium sulfide, prepn. of, by reaction of
```

ruthenium carbonyl and sulfur) TI12068-85-8, Iron disulfide (deposition of film of, by reaction of iron carbonyl and sulfur and hydrogen sulfide and thiourea) 1317-33-5P, Molybdenum disulfide, preparation IT(deposition of film of, by reaction of molybdenum carbonyl and sulfur) IT12138-09-9, Tungsten disulfide (deposition of film of, by reaction of tungsten carbonyl and sulfur) ANSWER 10 OF 27 HCA COPYRIGHT 2003 ACS on STN L55 108:153728 Cathodes for photochargeable solid-state Kanbara, Teruhisa; Tonomura, Tadashi; Kondo, battery. Shigeo (Dodensei Muki Kagobutsu Gijutsu Kenkyu Kumiai, Japan). Kokai Tokkyo Koho JP 63004558 A2 19880109 Showa, 5 (Japanese). CODEN: JKXXAF. APPLICATION: JP 1986-148780 19860625. The photochargeable battery consists of a Cu-based AB anode, a Cu+-conducting electrolyte, and an n-type WX2 Thus, a cathode layer of a 2:3 mixt. of WSe2-RbCu4I1.5Cl3.5 electrolyte (I); an electrolyte layer; and an anode layer of a 4:19:5 mixt. of Cu powder, Cul.59S, and I were pressed together to form a pellet, which was attached with a In-Sn oxide-coated glass plate cathode collector and a carbon fiber-SBR anode collector to form a battery having better charging-discharging performance at 60.degree. than a battery using a ZrS2-contg. cathode. 12067-46-8 12067-76-4 IT(cathodes, for photochargeable solid-state copper battery) 12067-46-8 HCA RN Tungsten selenide (WSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNSe W Se 12067-76-4 HCA RNTungsten telluride (WTe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNTe == W == Te IC ICM H01M004-58 ICS H01L031-04; H01M004-02; H01M010-36 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC tungsten selenide photochargeable battery cathode ST TIBatteries, secondary (copper/tungsten chalcogenide, solid-state, photochargeable) IT Cathodes (battery, tungsten chalcogenide, photochargeable)

12138-09-9

12067-46-8 12067-76-4

ΙT

(cathodes, for photochargeable solid-state copper battery) HCA\ COPYRAGHT 2003 ACS on STN L55 ANSWER AT OF 27 HCA COPINIGHI 2003 ACS ON SIL.

108:153727 Photochargeable secondary solid-state battery.

Kanbara Teruhisa; Tonomura, Tadashi; Kondo, Shigeo (Dodensei Muki Kagobutsu Gijutsu Kenkyu Kumiai, Japan). Jpn. Kokai Tokkyo Koho JP L55 27 63004557 A2 19880109 Showa 5 (Japanese). CODEN: JKXXAF. APPLICATION: JP 1986-148779 \ 19860625. The photochargeable **battery** consists of a Cu-based **anode**, a Cu+-conducting solid electrolyte, and a AΒ (cathode) mainly consisting of n-type MoX2 (X = S, Se, or Te), and is charged by illuminating the cathode. Thus, a battery waas prepd. by pressing a cathode layer of a 2:3 (wt.) MoSe2-RbCu4I1.5Cl3.5 (I, electrolyte) mixt., an electrolyte layer; and an anode layer of a 4:19:5 (wt.) Cu powder-Cu1.59S-I mixt. to form a pellet; and connecting a cathode collector of an In-Sn oxide-coated glass and an anode collector of a carbon-SBR composite to the resp. electrodes. When cycled at 60.degree. by 1-h discharging at 100-.mu.A and 1-h charging by illuminating the cathode with a 100-W Xe lamp from a 50-cm distance, this battery had better performance than a battery using a ZrS2 cathode. IT12058-18-3 12058-20-7 (cathodes, for photo-chargeable solid-state batteries) RN12058-18-3 HCA CNMolybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) Se Mo Se RN12058-20-7 HCA CNMolybdenum telluride (MoTe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) Te--- Mo--- Te IC ICM H01M004-58 ICS H01L031-04; H01M004-02; H01M010-36 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) STmolybdenum selenide photocharqeable battery cathode Cathodes IT (battery, molybdenum chalcogenide, photochargeable) IT7440-05-3, Palladium, uses and miscellaneous 7440-06-4, Platinum, uses and miscellaneous (cathodes contg., molybdenum chalcogenide, for photochargeable solid-state battery) IT 1317-33-5, uses and miscellaneous 12058-18-3 12058-20-7

(cathodes, for photo-chargeable solid-state
batteries)

L55 ANSWER 12 OF 27 HCA COPYRIGHT 2003 ACS on STN 108:153645 Solar-powered secondary **batteries**. Kanbara, Teruhisa; Tonomura, Tadashi (Dodensei Muki Kagobutsu Gijutsu Kenkyu Kumiai, Japan). Jpn. Kokai Tokkyo Koho JP 62259359 A2 19871111 Showa, 4 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1986-102343 19860502.

The title batteries have a Cu anode, a Cu+-conducting solid electrolyte, and n-type ZrX2 (X = Se, Te) cathode. Thus, layers of 60 mg 2:3 (wt.)

ZrSe2-RbCu4I1.5Cl3.5 (I) mixt., 50 mg I; and 50 mg 4:19:5 (wt.) mixt. of Cu, Cu1.59S, and I were pressed to form a battery with the cathode covered by a glass plate having an In-Sn oxide transparent layer. The electrodes are connected by a diode to prevent reverse current during charging. When charged by a 100-W Xe lamp from a 50-cm distance and discharged at 100 .mu.A, it showed less output-voltage decrease than a battery using a ZrS2 cathode.

IT 113671-81-1 113671-82-2

(n-type cathodes, for solar-charged solid-state copper batteries)

RN 113671-81-1 HCA

CN Zirconium selenide (ZrSe1.8-2.1) (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number	
Se Zr	1.8 - 2.1	7782-49-2 7440-67-7	

RN 113671-82-2 HCA

CN Zirconium telluride (ZrTe1.8-2.1) (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
Te	1.8 - 2.1	13494-80-9
Zr	1	7440-67-7

IC ICM H01M010-46

ICS H01L031-04; H01M004-58; H01M010-36

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

ST solar charged secondary battery; zirconium selenide

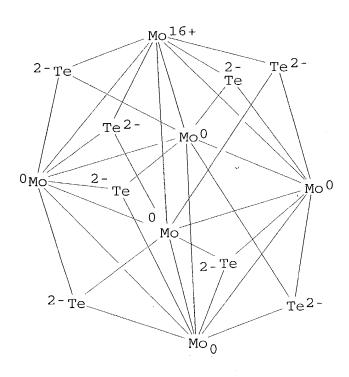
battery cathode
IT Cathodes

(battery, solar-charged, n-type zirconium selenide or telluride)

IT 113671-81-1 113671-82-2

(n-type cathodes, for solar-charged solid-state copper batteries)

```
L55
    ANSWER 13 OF 27 HCA COPYRIGHT 2003 ACS on STN
106:159511 Lithium electrochemical cells at low
    voltage: decomposition of molybdenum and tungsten dichalcogenides.
     Selwyn, L. S.; McKinnon, W. R.; Von Sacken, U.; Jones, C. A. (Div.
    Chem., Natl. Res. Counc. Canada, Ottawa, ON, K1A OR9, Can.). Solid
     State Ionics, 22(4), 337-44 (English) 1987. CODEN:
     SSIOD3. ISSN: 0167-2738.
AB
    Results are given for low voltage discharges of Li
    electrochem. cells contg. layered
    dichalcogenides of Mo and W. The reversible intercalation of Li at
    high voltages occurs only for phases with octahedrally coordinated
    metal atoms, but all phases decomp. irreversibly at low voltages to
    a mixt. of Mo or W and Li2X, where X is S, Se, or Te. Li can be
    removed electrochem. from these mixts. at a voltage that correlates
    with the free energies of formation of Li2X.
    12058-18-3, Molybdenum diselenide 12058-20-7,
ΙT
    Molybdenum ditelluride 12067-46-8, Tungsten diselenide
     57620-27-6
        (cathodes, in lithium batteries,
        intercalation in relation to)
    12058-18-3 HCA
RN
    Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
Se Mo Se
RN
    12058-20-7 HCA
CN
    Molybdenum telluride (MoTe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
Te≡ Mo≡ Te
    12067-46-8 HCA
RN
    Tungsten selenide (WSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
Se== W== Se
     57620-27-6 HCA
RN
    Molybdenum, octa-.mu.3-telluroxohexa-, octahedro (9CI)
                                                             (CA INDEX
CN
    NAME)
```



52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC

lithium intercalation tungsten molybdenum chalcogenide; ST

cathode tungsten molybdenum chalcogenide lithium

IΤ Cathodes

> (battery, dichalcogenides of molybdenum or tungsten, with lithium anodes)

TI7439-93-2, Lithium, uses and miscellaneous

> (anodes, in batteries with dichalcogenides of molybdenum or tungsten as cathodes)

1317-33-5, Molybdenum disulfide, uses and miscellaneous ΙT 12058-18-3, Molybdenum diselenide 12058-20-7, Molybdenum ditelluride 12067-46-8, Tungsten diselenide 12138-09-9, Tungsten disulfide 57620-25-4, Molybdenum sulfide (Mo6S8) 57620-27-6

> (cathodes, in lithium batteries, intercalation in relation to)

ANSWER 14 OF 27 HCA COPYRIGHT 2003 ACS on STN

104:171377 Rechargeable solid electrolyte cells with a copper ion conductor, Rb4Cu16I7-xCl13+x. Kanno, Ryoji; Takeda, Yasuo; Oda, Yasuhiro; Ikeda, Hiroyuki; Yamamoto, Osamu (Fac. Eng., Mie Univ., Tsu, 514, Japan). Solid State Ionics, 18-19(2), 1068-72 (English) 1986, CODEN: SSIOD3. ISSN: 0167-2738.

A high performance secondary solid electrolyte battery AΒ with a high Cu ion conductor, Rb4Cu16I6.8Cl13.2, was developed. cell using Cu-Chevrel phase Cu4Mo6S8 as an anode and a layered Cu intercalation compd. NbS2 as a cathode, had a high discharge capacity and good rechargeability. At a c.d. IT

RN

CN

Te-Ta-Te

of 150 .mu.A/cm2, the cathode capacity was 60 mA-h/g (0.3 electron/NbS2) and at 750 .mu.A/cm2 it was 48 mA-h/g (0.24 electron/NbS2). On the charge-discharge cycle test of 75 .mu.A/cm2 and 0.3% electron/NbS2 depth, the cell did not show a significant deterioration in the charge-discharge curves during .gtoreg.500 12034-77-4 12034-83-2 12039-55-3 12058-18-3 12058-20-7 12067-45-7 12067-66-2 12067-76-4 (cathodes, battery, with copper rubidium chloride iodide electrolyte, performance of) 12034-77-4 HCA Niobium selenide (NbSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) Se Nb Se 12034-83-2 HCA Niobium telluride (NbTe2) (7CI, 8CI, 9CI) (CA INDEX NAME) Te Nb Te 12039-55-3 HCA Tantalum selenide (TaSe2) (7CI, 8CI, 9CI) (CA INDEX NAME) Se Ta Se 12058-18-3 HCA Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) Se Mo Se 12058-20-7 HCA Molybdenum telluride (MoTe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) Te--- Mo--- Te 12067-45-7 HCA Titanium selenide (TiSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) Se Ti Se 12067-66-2 HCA Tantalum telluride (TaTe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

```
RN
     12067-76-4 HCA
CN
     Tungsten telluride (WTe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
Te == W == Te
CC
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
     Section cross-reference(s): 72
ST
     battery solid electrolyte copper conductor
IT
     Batteries, secondary
        (copper molybdenum sulfide-metal chalcogenide, with copper
        rubidium chloride iodide electrolyte, performance of)
IT
     58051-93-7
        (anodes, battery, with copper rubidium
        chloride iodide electrolyte, performance of)
     1317-33-5, uses and miscellaneous 12034-77-4
IT
                  12039-13-3
                               12039-15-5 12039-55-3
     12034-83-2
     12058-18-3 12058-20-7 12067-45-7
     12067-66-2 12067-76-4
                             12138-09-9
                                          12143-72-5
     12164-84-0
                  12166-28-8
                             39290-91-0
                                            42821-47-6 42821-48-7
     52226-00-3
        (cathodes, battery, with copper rubidium
        chloride iodide electrolyte, performance of)
IT
     56188-83-1D, solid solns. with copper rubidium chloride
     63310-92-9D, solid solns. with copper rubidium iodide
        (electrolytes, copper molybdenum sulfide-niobium sulfide
        battery, performance of)
     ANSWER 15 OF 27 HCA COPYRIGHT 2003 ACS on STN
L55
102:223362 Progress in the SERI-DOE photoelectrochemical cell program.
     Wallace, William (Sol. Energy Res. Inst., Golden, CO, 80401, USA).
     Conference Record of the IEEE Photovoltaic Specialists Conference,
     16th, 1066-71 (English) 1982. CODEN: CRCNDP.
                                                    ISSN:
     0160-8371.
AΒ
     The concept of photoelectrochem. storage was investigated and
     demonstrated in studies involving 3-electrode in-situ and
     4-electrode redox storage cells. For these cells system
     efficiencies of 1-3% were obtained to date for the net conversion of
     light to elec. energy. SERI also monitors the tech. progress in the
     Texas Instruments solar energy system program which involves
     conversion of light into elec. and thermal energy in a system which
     incorporates electrochem. storage. Research on polycryst. thin-
     film n-CdSe and n-CdSexTe1-x based photoelectrochem. cells
     resulted in achievement of .ltoreq.7% efficiencies for the direct
     conversion of light into elec. energy without
     storage in devices contg. a sulfide/polysulfide electrolyte.
     Higher efficiencies are possible with improved polycryst. thin
     films and alternate electrolytes.
IT
     1306-25-8D, solid solns. with cadmium selenide
        (electrodes, photoelectrochem. cells based on thin-
```

film cadmium selenide and, progress in)

```
RN
     1306-25-8 HCA
     Cadmium telluride (CdTe) (9CI) (CA INDEX NAME)
CN
Cd== Te
     1306-24-7, uses and miscellaneous
IT
        (electrodes, photoelectrochem. cells based on thin-
        film cadmium selenide telluride and, progress in)
RN
     1306-24-7 HCA
     Cadmium selenide (CdSe) (9CI) (CA INDEX NAME)
CN
Cd == Se
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
     cadmium selenide telluride photoelectrochem cell; photoelectrochem
ST
     cell development; redox photoelectrochem cell development; solar
     energy conversion storage
IT
     Energy
        (solar, conversion and storage of, progress in)
     1306-24-7D, solid solns. with cadmium telluride 1306-25-8D
IT
     , solid solns. with cadmium selenide
        (electrodes, photoelectrochem. cells based on thin-
        film cadmium selenide and, progress in)
IT
     1306-24-7, uses and miscellaneous
        (electrodes, photoelectrochem. cells based on thin-
        film cadmium selenide telluride and, progress in)
     ANSWER 16 OF 27 HCA COPYRIGHT 2003 ACS on STN
102:206554 Applications of electrogenerated conducting polymers in
     electrochemical photovoltaic cells. Noufi, Rommel
     (Sol. Electr. Convers. Res. Div., Sol. Energy Res. Inst., Golden,
     CO, 80401, USA). Conference Record of the IEEE Photovoltaic
     Specialists Conference, 16th, 1293-8 (English) 1982.
     CODEN: CRCNDP. ISSN: 0160-8371.
AB
     The electrochem. generated conducting polypyrrole PP [30604-81-0]
     films protect n-type semiconductor photoelectrodes from
     degrdn. while permitting electron exchange between the semiconductor
     and the electrolyte. The performance characteristics and stability
     of PP-covered GaAs, Si, CdTe, CdSe, and CdS photoelectrodes are
     discussed.
     1306-24-7, uses and miscellaneous 1306-25-8, uses
TΤ
     and miscellaneous
        (electrodes from polypyrrole-coated,
        photoelectrochem., performance and stability of)
     1306-24-7 HCA
RN
     Cadmium selenide (CdSe) (9CI) (CA INDEX NAME)
CN
```

Cd Se

```
RN
     1306-25-8 HCA
     Cadmium telluride (CdTe) (9CI) (CA INDEX NAME)
CN
Cd==Te
CC
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
     Section cross-reference(s): 38, 72
ST
     polypyrrole coating photoelectrochem electrode;
     silicon polypyrrole photoelectrochem electrode; sulfide
     cadmium polypyrrole photoelectrochem electrode; selenide
     cadmium polypyrrole photoelectrochem electrode; gallium
     arsenide polypyrrole photoelectrochem electrode; cadmium
     telluride polypyrrole photoelectrochem electrode
IT
        (photoelectrochem., polypyrrole-coated, performance and
        stability of)
     30604-81-0
IT
        (electrodes coated with, photoelectrochem.,
        performance and stability of)
IT
     1303-00-0, uses and miscellaneous
                                         1306-23-6, uses and
     miscellaneous 1306-24-7, uses and miscellaneous
     1306-25-8, uses and miscellaneous 7440-21-3, uses and
     miscellaneous
        (electrodes from polypyrrole-coated,
        photoelectrochem., performance and stability of)
                     HCA COPYRIGHT 2003 ACS on STN
     ANSWER 17 OF 27
102:206541 II-VI thin film electrochemical
     photovoltaic cells. Russak, Michael A.; Reichman, Joseph
     (Res. Dev. Cent., Grumman Aerosp. Corp., Bethpage, NY, 11714, USA).
     Conference Record of the IEEE Photovoltaic Specialists Conference,
     16th, 1057-61 (English) 1982. CODEN: CRCNDP.
     0160-8371.
     Group 12-16 (IIB-VIA) compd. thin film and thin-
AB
     film heterostructure electrodes for use in
     photoelectrochem. cells were produced from their constituent
     elements using a 3-source vacuum-evapn. system. CdSe films
     with a wide variation in electronic properties were produced,
     characterized, and evaluated with efficiencies of .ltoreq.6.5% being
     recorded for films <2.0.mu. thick. ZnSe/CdSe
     heterostructure produced in situ yielded cells with open-circuit
     voltages >0.6 V and efficiencies >5.5%. Simultaneous elemental
     evapn. was also used to make CdSe1-xTex (x = 0.05-0.40) thin
     films, which had efficiencies of .ltoreq.7%. Also, the use
     of an aq. ferro-ferricyanide electrolyte resulted in efficiencies
     >7% for CdSe thin films and .apprx.15% for CdSe
     single-crystal electrodes.
```

(electrodes from cadmium selenide covered with,

photoelectrochem., prepn. and properties of)
RN 1315-09-9 HCA

1315-09-9P

IT

CNZinc selenide (ZnSe) (9CI) (CA INDEX NAME) Se== Zn ΙT 1306-24-7P, uses and miscellaneous 1306-25-8DP, solid solns. with cadmium selenide (electrodes, photoelectrochem., prepn. and properties of) 1306-24-7 HCA RN Cadmium selenide (CdSe) (9CI) CN(CA INDEX NAME) Cd Se 1306-25-8 HCA RNCNCadmium telluride (CdTe) (9CI) (CA INDEX NAME) Cd== Te CC52-2 (Electrochemical, Radiational, and Thermal Energy Technology) ST cadmium selenide photoelectrochem electrode; zinc selenide photoelectrochem electrode; telluride selenide cadmium photoelectrochem electrode; ferrocyanide selenide photoelectrochem cell IT Photoelectric devices, solar (photoelectrochem., Group 12-16 (IIB-VIA) compd. thinfilm) IT 1315-09-9P (electrodes from cadmium selenide covered with, photoelectrochem., prepn. and properties of) IT1306-24-7DP, solid solns. with cadmium telluride 1306-24-7P , uses and miscellaneous 1306-25-8DP, solid solns. with cadmium selenide (electrodes, photoelectrochem., prepn. and properties of) ANSWER 18 OF 27 HCA COPYRIGHT 2003 ACS on STN 100:37151 Photoelectrochemical electrodes. Williams, Roger M.; Rembaum, Alan (United States National Aeronautics and Space Administration, USA). U.S. Pat. Appl. US 376306 A0 19830930, 29 pp. Avail. NTIS Order No. PAT-APPL-6-376 306. (English). CODEN: XAXXAV. APPLICATION: US 1982-376306 19820510. The surface of a moderate band-gap semiconductor such as p-type MoS2 AB is modified to contain an adherent film of charge-mediating ionene polymer contg. an electroactive unit such as bipyridinium. Electron transport between the electrode and the mediator film is favorable and photocorrosion and recombination processes are suppressed. Incorporation of particles

of catalyst such as Pt within the **film** provides a redn. in overvoltage. The polymer **film** is readily deposited on the

electrode surface and can be rendered stable by ionic or addn. crosslinking. Catalyst can be predispersed in the polymer film or a salt can be impregnated into the film Thus natural p-MoS2 crystals showed greatly and reduced there. enhanced visible light to chem. energy conversion efficiency following modification with a film composed of a charge-mediating polymer (crosslinked 4,4'-bipyridylchloromethylstyrene copolymer [88379-78-6]) and catalyst (Pt). Onset of dark currents for H synthesis indicated that the dark overpotential is reduced to .apprx.100 mV. Photoelectrochem. energy conversion efficiency was increased from .apprx.0 to 3.9% on modifying MoS2 surface with 2.5 mg polymer/cm2 and 0.25 mg Pt/cm2. 12058-18-3 (electrodes, coated with polymer contg. dispersed platinum, photoelectrochem.-cell) 12058-18-3 HCA Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) Se Mo Se 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 38, 72 photoelectrochem cell electrode molybdenum sulfide; bipyridyl chloromethylstyrene polymer electrode photoelectrochem; platinum bipyridyl chloromethylstyrene polymer electrode; hydrogen manuf photoelectrochem cell (photoelectrochem., molybdenum chalcogenide, coated with ionic polymer contg. dispersed catalyst) 88379-78-6 (crosslinked, electrodes from semiconductor coated with layer of catalysts-contg., photoelectrochem.-cell) 7440-06-4, uses and miscellaneous (electrodes from semiconductor coated with polymer contg. dispersed, photoelectrochem.-cell) 1317-33-5, uses and miscellaneous 12058-18-3 (electrodes, coated with polymer contg. dispersed platinum, photoelectrochem.-cell) 88375-42-2 88375-43-3 88375-41-1 88375-40-0 32168-10-8 88375-45-5 88375-44-4 (ionene, electrodes from semiconductor coated with layer of catalysts-contg., photoelectrochem.-cell) 7782-44-7P, preparation 1333-74-0P, preparation (manuf. of, in photoelectrochem. cells, electrodes coated with ionic polymert contg. dispersed catalyst for) ANSWER 19 OF 27 HCA COPYRIGHT 2003 ACS on STN

98:110704 Semiconductors and semiconductor photoelectrodes.

Joost; Cahen, David; Hodes, Gary (Yeda Research and Development Co.,

IT

RN

CN

CC

ST

IT

IT

IT

IT

IT

ΙT

```
Ltd., Israel). U.S. US 4368216 A 19830111, 7 pp.
     Cont.-in-part of U.S. 4,296,188. (English). CODEN: USXXAM.
     APPLICATION: US 1980-173814 19800730. PRIORITY: IL 1979-58003
     19790808; US 1980-134665 19800327.
     Photoelectrodes are manufd. by prepg. a slurry of .gtoreq.1
AB
     semiconductor, a flux, and a liq. vehicle, applying a layer
     of the slurry to an elec. conductive substrate, and annealing the
             Thus, powd. CdSe and ZnCl2 were ground together with
     a mixt. of 5% nonionic detergent in water. This paint was applied
     to a piece of preheated Ti. The coated Ti was heated at
     650.degree. for 12 min in Ar contg. 20 ppm O. A photoelectrochem.
     cell contq. this electrode, a sulfided brass gauze counter
     electrode, and an electrolyte of aq. 1M KOH, 1M Na2S.9H2O,
     and 1M S gave under air-mass-1 conditions a short-circuit current
     (I) of 26.2 mA, an open-circuit voltage (V) of 530 mV, and a
     photopotential (P) of 388 mV over an optimal load of 24 .OMEGA...
     After etching the CdSe electrode in 3% HNO3 in concd. HCl
     the same photoelectrochem. cell yielded I = 36.5 mA, V = 605 mV, and
     P = 424 \text{ mV} over an optimal load of 17 .OMEGA.. After the
     electrode was dipped for 3 s in 1M aq. ZnCl2, the cell
     yielded I = 36.8 \text{ mA}, V = 660 \text{ mV}, and P = 463 \text{ mV} over an optimal load
     of 18 .OMEGA..
IT
     1306-24-7P, uses and miscellaneous 1306-25-8DP,
     solid solns. with cadmium sulfide and/or cadmium selenide
     1306-25-8P, uses and miscellaneous 1315-09-9P
     1315-11-3P
        (electrodes, photoelectrochem.-cell, prepn. and
        performance of)
     1306-24-7
               HCA
RN
     Cadmium selenide (CdSe) (9CI) (CA INDEX NAME)
CN
Cd Se
     1306-25-8 HCA
RN
     Cadmium telluride (CdTe) (9CI) (CA INDEX NAME)
CN
Cd = Te
RN
     1306-25-8
                HCA
     Cadmium telluride (CdTe) (9CI) (CA INDEX NAME)
CN
Cd=Te
RN
     1315-09-9 HCA
     Zinc selenide (ZnSe) (9CI) (CA INDEX NAME)
CN
```

Se = Zn

```
RN
     1315-11-3 HCA
     Zinc telluride (ZnTe) (8CI, 9CI) (CA INDEX NAME)
CN
Te = Zn
     B05D003-02; H01M006-36
IC
NCL
     427074000
CC
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
     semiconductor photoelectrochem cell electrode; cadmium
ST
     selenide photoelectrochem cell electrode; zinc chloride
     flux photoelectrochem electrode; sodium sulfide
     photoelectrochem cell electrolyte
IT
     Electrodes
        (photoelectrochem., semiconductive, prepn. and performance of)
     7440-47-3, uses and miscellaneous
IT
        (electrodes from semiconducting layer on
        steel plated with, photoelectrochem.-cell)
     7440-32-6, uses and miscellaneous 7782-42-5, uses and
IT
     miscellaneous
        (electrodes from semiconducting layer on,
        photoelectrochem.-cell)
     1303-00-0P, uses and miscellaneous 1306-23-6DP, solid solns. with
IT
     cadmium selenide and/or cadmium telluride 1306-23-6P, uses and
                    1306-24-7DP, solid solns. with cadmium sulfide
     miscellaneous
     and/or cadmium telluride 1306-24-7P, uses and
     miscellaneous 1306-25-8DP, solid solns. with cadmium
     sulfide and/or cadmium selenide 1306-25-8P, uses and
                     1315-09-9DP, solid solns. with cadmium selenide
     miscellaneous
     1315-09-9P 1315-11-3P
                             1317-33-5P, uses and
     miscellaneous
                     12018-94-9P
        (electrodes, photoelectrochem.-cell, prepn. and
        performance of)
IT
     7699-45-8
        (flux from cadmium sulfate and; in manuf. of semiconductor
        electrodes for photoelectrochem. cells)
IT
     10124-36-4
        (flux from zinc chloride and, in manuf. of semiconductor
        electrodes for photoelectrochem. cells)
IT
     10108-64-2
        (flux, in manuf. of semiconductor electrodes for
        photoelectrochem. cells)
     1313-82-2, uses and miscellaneous
IT
        (photoelectrochem.-cell electrolyte,
        semiconductor)
     ANSWER 20 OF 27 HCA COPYRIGHT 2003 ACS on STN
97:117301 Electrochemical photovoltaic cells
     /stabilization and optimization of II-VI semiconductors. Noufi, R.;
     Tench, D.; Warren, L. (Rockwell Int. Corp., Thousand Oaks, CA, USA).
```

Report, SERI/TR-9276-T3; Order No. DE82002093, 55 pp. Avail. NTIS From: Energy Res. Abstr. 1982, 7(5), Abstr. No. 10440 (English)

1981.

The goal of this program is to provide a basis for designing a AΒ practical electrochem. solar cell based on the Group II-VI compd. semiconductors. Emphasis is on developing new electrolyte redox systems and electrode surface modifications which will stabilize the Group II-VI compds. against photodissoln. without seriously degrading the long-term solar response. Although the MeOH ferro-ferricyanide soln. itself was photolytically unstable, study of this system led to the identification of more promising nonaq. redox electrolytes. work on redox couple stabilization of n-type CdX photoanodes was focused on both 1- and 2-electron couples. Very promising results were obtained for the alkylammonium chloroferrate (II, III) couple in Conducting polymer films of polypyrrole photoelectrochem. deposited onto n-type semiconductors were shown to protect these electrode materials from photodecompn. while permitting electron exchange with the electrolyte. In a basic aq. ferro-ferricyanide electrolyte contg. cyanide ion, the measured open-circuit voltage for n-type CdTe was 1.3 V, which is practically the bandgap for this material. It now appears that polypyrrole films are to some extent permeable to solvent/solute species since the film stability depends on the nature of the redox electrolyte.

IT 1306-24-7, uses and miscellaneous

(anodes, photoelectrochem., stabilization of)

RN 1306-24-7 HCA

CN Cadmium selenide (CdSe) (9CI) (CA INDEX NAME)

Cd = Se

IT **1306-25-8**, properties

(elec. open-circuit potential of, in basic aq. soln. contg. cyanoferrate couple and cyanide ion)

RN 1306-25-8 HCA

CN Cadmium telluride (CdTe) (9CI) (CA INDEX NAME)

Cd = Te

CC 72-2 (Electrochemistry)

Section cross-reference(s): 52, 76

solar cell semiconductor stabilization optimization; Group II chalcogenide photoelectrode stabilization; photoanode cadmium selenide stabilization; polypyrrole film stabilization photoelectrode; polyaniline film stabilization photoelectrode

IT Electrodes

(photoelectrochem., Group II chalcogenides, stabilization of)

IT Anodes

(photoelectrochem., cadmium selenide, stabilization of)

IT 1306-24-7, uses and miscellaneous

(anodes, photoelectrochem., stabilization of) 30604-81-0 ΙT (coatings of, on Group IIV chalcogenides electrodes, quality compn. prevention in relation to) 1306-25-8, properties IT(elec. open-circuit potential of, in basic aq. soln. contg. cyanoferrate couple and cyanide ion) 25233-30-1 IT(films, in stabilization of Group II chalcogenide semiconductor electrons) ANSWER 21 OF 27 HCA COPYRIGHT 2003 ACS on STN 94:211521 Electrochemical photovoltaic cells /stabilization and optimization of II-VI semiconductors. Noufi, R.; Tench, D.; Warren, L. (Rockwell Int. Corp., Thousand Oaks, CA, USA). Report, SERI/TR-8002/T1, 73 pp. Avail. NTIS From: Energy Res. Abstr. 1980, 5(24), Abstr. No. 37851 (English) 1980. Stabilization of n-CdSe against photodissoln. was achieved for the ΆB MeOH-(Et4N)4Fe(CN)6-(Et4N)3Fe(CN)6 system. No degrdn. of the photocurrent or the electrode surface, even in the presence of traces of H2O, was obsd. for runs of .ltoreq.700 h at 6 mA/cm2 and approx. air-mass-1 intensity. With higher quality single-crystal CdSe, stable short-circuit photocurrents of 15-17 mA/cm2 and an open-circuit voltage of 0.7 V were obtained, corresponding to a conversion efficiency of .apprx.5%. Preliminary evaluation of a series of S-contg. 1,2-dithiolene metal complexes for stabilization of Cd chalcogenide photoanodes in soln. was completed. For the 1st time, a conducting polymer film (derived from pyrrole) was electrochem. deposited on a semiconductor This could represent a breakthrough in the use electrode. of hydrophobic films to protect semiconductor photoanodes from dissoln./degrdn. Mixed CdSe-CdTe solid soln. electrodes were found to exhibit a min. in both the flatband potential and the bandgap at .apprx.65% CdTe. Both of these shifts would have a detrimental effect on the solar conversion efficiency. 1306-24-7, uses and miscellaneous IT(anodes, photoelectrochem.-cell, optimization and stabilization of) RN1306-24-7 HCA Cadmium selenide (CdSe) (9CI) (CA INDEX NAME) CNCd==Se 1306-25-8D, solid solns. with cadmium selenide IT(anodes, photoelectrochem.-cell, properties of) 1306-25-8 HCA RN Cadmium telluride (CdTe) (9CI) (CA INDEX NAME) CN

Cd==Te

```
52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
     1306-24-7, uses and miscellaneous
TI
         (anodes, photoelectrochem.-cell, optimization and
        stabilization of)
     1306-24-7D, solid solns. with cadmium telluride 1306-25-8D
IT
     , solid solns. with cadmium selenide
         (anodes, photoelectrochem.-cell, properties of)
     ANSWER 22 OF 27 HCA COPYRIGHT 2003 ACS on STN
L55
94:106551 Active electrodes for electrochemical
     battery cells with redox systems. Hodes, Gary;
     Manassen, Joost; Cahen, David (Yeda Research and Development Column Ltd., Israel). Ger. Offen, DE 3004262 19800904, 20 pp.
     (German). CODEN: GWXXBX. APPLICATION DE 1980-3004262 19800206.
     Electrodes for the title S/S2-, Se/Se2-, and Te/Te2-
AB
     batteries and photoelectrochem. cells comprise an active
     material-coated substrate of steel, stainless'steel,
     porous C, graphite, Co, Ti, Ta, W, Mo, V, and Cr. The possible active materials are CoS, Cu2S, RuS2, MoS2, PbSe, Cu2Se, and NiTe.
     Thus, several chalcogenide-coated stainless steel
     electrodes were prepd. and their properties in
     photoelectrochem. cells were measured.
     1314-05-2 1314-91-6D, solid solns. with lead
IT
     selenide 11115-77-8 12017-13-9D, solid solns.
     with cobalt selenide 12142-88-0D, solid solns. with nickel
     selenide 39280-96-1
         (electrodes from stainless steel coated with,
         photoelectrochem.-cell)
     1314-05-2 HCA
RN
     Nickel selenide (NiSe) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
Ni == Se
     1314-91-6 HCA
RN
     Lead telluride (PbTe) (6CI, 8CI, 9CI) (CA INDEX NAME)
CN
Pb== Te
     11115-77-8 HCA
RN
     Cobalt telluride (9CI) (CA INDEX NAME)
CN
                                             Component
  Component
                        Ratio
                                          Registry Number
                                               13494-80-9
Te
                          Х
                                               7440-48-4
                          Х
Co
```

Cobalt telluride (CoTe) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

12017-13-9 HCA

RN

CN

```
Co== Te
```

12142-88-0 HCA RNNickel telluride (NiTe) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CN

Ni = Te

39280-96-1 HCA RN

39280-96-1

CN	N Lead telluride (9CI) (CA INDEX NAME)			
	mponent	Ratio	Component Registry Number	
Te Pb		x x	13494-80-9 7439-92-1	
IC H01M004-58; H01M010-36; H01M014-00 CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) ST chalcogenide electrode photoelectrochem cell; stainless steel chalcogenide coating electrode; battery redox chalcogenide electrode; sulfide electrode photoelectrochem cell; selenide electrode photoelectrochem cell; telluride electrode photoelectrochem cell				
IT	•			
IT	T 22205-45-4 (electrodes from brass coated with, photoelectrochemcell)			
IT	T 12597-68-1, uses and miscellaneous (electrodes from chalcogenide-coated, photoelectrochemcell)			
ΙΤ	12597-71-6, uses and miscellaneous (electrodes from copper sulfide-coated, photoelectrochemcell)			
IT	<pre>1317-33-5, uses and miscellaneous (electrodes from molybdenum coated with, photoelectrochemcell)</pre>			
IT				
IT	1314-05-21 solns. wit lead seler 12017-13-9	o, solid solns. with a ch lead selenide 1314 nide 1317-42-6 11 op, solid solns. with op, solid solns. with op, solid solns. with op, solid solns. with		

(electrodes from stainless steel coated with, photoelectrochem.-cell)

ANSWER 23 OF 27 HCA COPYRIGHT 2003 ACS on STN 91:143244 On the photopotential output of electrochemical solar cells based on layer-type d-band semiconductors. Tributsch, H.; Gerischer, H.; Clemen, C.; Bucher, E. (Fritz-Haber-Inst., Max-Planck-Ges., Berlin, D-1000/33, Fed. Rep. Ger.). Berichte der Bunsen-Gesellschaft, 83(7), 655-8 (English) 1979. CODEN: BBPCAX. ISSN: 0005-9021. A comparative study of Mo and W-dichalcogenides in contact with AΒ various redox electrolytes revealed that in abs. values as well as in relation to the energy gap, n-type WSe2 is producing the largest photopotential output. It exceeds 0.55 V in presence of several redox couples (Fe2+/3+, hydroquinone/quinone pH = 10, Fe(CN)63-/4-, Ru3+/4+, Br-/Br2, Ce3+/4+) and amts. to more than 0.7 V in presence of I-/I2. Addn. of small quantities of iodide can increase the photopotential output in presence of redox systems with lower redox potential by .ltoreq.0.1 V. This gives further evidence for the specific photochem. surface activity of iodide on layer -type electrodes and is an example for the influence of surface states on the efficiency of energy conversion. The higher photopotential output of WSe2 as compared to that of MoSe2 (max. value I-/I2, 0.56 V) is explained in terms of a neg. shift of the conduction band edge. With respect to the efficiency of electrochem. solar energy conversion WSe2 is considered to be the most promising compd. among layer-type d-band semiconductors. 12058-18-3 12058-20-7 12067-46-8 IT (electrodes, photoelectrochem. cells contg., properties of) RN12058-18-3 HCA Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNSe Mo Se 12058-20-7 HCA RNMolybdenum telluride (MoTe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNTe--- Mo--- Te RN 12067-46-8 HCA Tungsten selenide (WSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNSe== W== Se 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC

selenide tungsten photoelectrochem cell; molybdenum selenide

photoelectrochem cell; chalcogenide layer type

ST

```
photoelectrochem cell
     Iodides, uses and miscellaneous
IT
        (photoelectrochem. cells contg., properties of layer
        -type d-band semiconductors in)
     1317-33-5, uses and miscellaneous 12058-18-3 12058-20-7 12067-46-8
IT
        (electrodes, photoelectrochem. cells contg., properties
        of)
     ANSWER 24 OF 27 HCA COPYRIGHT 2003 ACS on STN
L55
         Electrode for electrolytic process involving hydrogen
     generation. Westerlund, H. Benny (Gow Enterprises Ltd., Can.).
     U.S. US 4089771 19780516, 6 pp. (English). CODEN:
              APPLICATION: US 1977-806409 19770614.
     USXXAM.
     A Ti bipolar electrode is described.
                                            The central core is
AΒ
     of Ti expanded metal sheet and an extension of this provides the
     anodic surface which may be coated, e.g., by Pt as is a well
     known practice. The cathodic surface may be activated to
     provide a hydride surface or coated with MoS2 or with Ag then MoS2.
     The electrode space is on the exposed side of the
     cathodic element and is of a non-conductive material such as
                        The electrode was tested in brine
     Kel-F or Teflon.
     electrolysis and ClO3- cells.
     1317-33-5, uses and miscellaneous
IT
        (coatings, on titanium porous bipolar electrodes)
     1317-33-5 HCA
RN
     Molybdenum sulfide (MoS2) (8CI, 9CI) (CA INDEX NAME)
CN
S = M_0 = S
     C25B011-03
IC
     204284000
NCL
     72-10 (Electrochemistry)
CC
     titanium bipolar electrode brine chlorate; molybdenum
ST
     sulfide coating titanium electrode
     Brines
IT
         (electrolysis of, porous bipolar titanium electrodes
        for)
     Coating materials
IT
         (molybdenum sulfide, on porous bipolar titanium
        electrodes)
     1317-33-5, uses and miscellaneous
IT
         (coatings, on titanium porous bipolar electrodes)
     7440-32-6, uses and miscellaneous
IT
         (electrodes, bipolar, porous, molybdenum disulfide
        coated)
     1333-74-0P, preparation
IT
         (generation of, electrode for)
IT
     14866-68-3P
         (manuf. of, porous bipolar titanium electrodes for)
```

L55 ANSWER 25 OF 27 HCA COPYRIGHT 2003 ACS on STN
89:46261 Chalcogenides of arsenic, antimony and bismuth as positive
electrodes in lithium batteries. Besenhard,
Juergen O. (Anorg.-Chem. Inst., Tech. Univ. Muenchen, Munich, Fed.
Rep. Ger.). Zeitschrift fuer Naturforschung, Teil B: Anorganische
Chemie, Organische Chemie, 33B(3), 279-83 (German) 1978.
CODEN: ZNBAD2. ISSN: 0340-5087.

AB Redox properties of layer-type chalcogenides of Group VA elements were investigated in molten salt and org. Li+-contg. electrolytes. Even at room temp., Bi2S3, Bi2Se3, and Bi2Te3 can be reduced with .apprx.100% efficiency to Li3Bi as final product. The redn. of Bi2S3 to Bi0 and LiS can be reversed in org. electrolytes.

IT 1304-82-1 12068-69-8

(cathodes, in batteries with lithium
anode, properties of)

RN 1304-82-1 HCA

CN Bismuth telluride (Bi2Te3) (8CI, 9CI) (CA INDEX NAME)

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

RN 12068-69-8 HCA

CN Bismuth selenide (Bi2Se3) (6CI, 8CI, 9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
=======================================		+====================================
Se	3	7782-49-2
Bi	2	7440-69-9

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

antimony chalcogenide battery cathode; arsenic chalcogenide battery cathode; bismuth chalcogenide battery cathode; sulfide bismuth battery cathode; selenide bismuth battery cathode; telluride bismuth battery cathode; lithium battery chalcogenide cathode

IT Cathodes

(battery, chalcogenides of arsenic, antimony and bismuth, properties of)

IT 1303-33-9 **1304-82-1** 1345-04-6 1345-07-9

12068-69-8

(cathodes, in batteries with lithium anode, properties of)

L55 ANSWER 26 OF 27 HCA COPYRIGHT 2003 ACS on STN
85:180084 Semiconductor-electrolyte photovoltaic energy
converter. Anderson, William W.; Anderson, Larry B. (Dep.
Electr. Eng., Ohio State Univ., Columbus, OH, USA). NASA Contract.
Rep., NASA-CR-143107,f., 1st, 1975, 702-12 (English) 1975.
CODEN: NSCRAQ.

AB Feasibility and practicality of a solar cell consisting of a semiconductor surface in contact with an electrolyte are considered. Characteristics of single crystal CdS, CdSe, CdTe, GaAs, and thinfilm CdS in contact with aq. and MeOH electrolytes are

reported.

IT 1306-24-7 1306-25-8

(electrodes, photoelectrochem.-cell, characteristics and soln. stability of)

RN 1306-24-7 HCA

CN Cadmium selenide (CdSe) (9CI) (CA INDEX NAME)

Cd== Se

RN 1306-25-8 HCA

CN Cadmium telluride (CdTe) (9CI) (CA INDEX NAME)

Cd== Te

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT Photoelectric cells

(solar, electrolyte-semiconductor, characteristics and practicality of)

1303-00-0, uses and miscellaneous 1306-23-6, uses and miscellaneous 1306-24-7 1306-25-8

(electrodes, photoelectrochem.-cell, characteristics and soln. stability of)

L55 ANSWER 27 OF 27 HCA COPYRIGHT 2003 ACS on STN
79:99856 Alkali metal/sulfur battery having a cathodic current collector coated with molybdenum disulfide. Strangham, Robert R.; Taplin, William H. (Dow Chemical Co.). (U.S. US 3749603 19730731, 6 pp. (English). CODEN: USXXAM. APPLICATION: USXXAM. 1972-244410 19720417.

An alkali metal/S battery having as a cathodic AΒ current collector a shaped member comprising a substrate on which there is a 50-5000 .ANG. thick coating of MoS2 is disclosed. substrate can be an outer layer or may constitute the entire member and consists essentially of Al, Mg or alloys contg. at least minor amts. of these metals. The elec. resistance of the coated cathode generally drops to a min. upon being contacted with a molten alkali metal polysulfide for a brief time. Test specimens were prepd. by cutting in half a 5/16 .times. 3 in. strip of the sheet or foil. After pretreatment, specimens were bolted to stainless steel lead-in wires, spaced apart by a 1/8 in. thick glass bar at their upper ends and inserted in a test tube. The tube was filled with Na2S4 in a dry box and placed in a heating block while connected to a low pressure purge of dry N. When the sulfide melted, the position of the strips was adjusted, if necessary, so that 10 cm2 of surface on each strip was in contact with the melt. A temp. of 300.degree. was maintained. A test voltage of 100 mV d.c. was applied, with the polarity being reversed automatically at 0.5-hr. intervals. The surface resistance after any interval of test duration was detd.

IT 1317-33-5

(coatings, on aluminum alloys, for cathodes for alkali metal-sulfur batteries) 1317-33-5 HCA RNMolybdenum sulfide (MoS2) (8CI, 9CI) (CA INDEX NAME) CN $S = M_0 = S$ IC H01M 136006000 NCL 77-2 (Electrochemistry) CC alkali metal sulfur battery; molybdenum sulfide ST cathode collector Batteries, secondary IT(alkali metal-sulfur, molybdenum disulfide coatings on cathode substrates in) Coating materials IT(molybdenum disulfide, on aluminum alloys, for cathodes for alkali metal-sulfur batteries) 12608-67-2 42614-09-5 IT (coatings on, of molybdenum disulfide, for cathode in alkali metal-sulfur batteries) 1317-33-5 IT(coatings, on aluminum alloys, for cathodes for alkali metal-sulfur batteries) => d 156 1-18 cbib abs hitstr hitind ANSWER 1 OF 18 HCA COPYRIGHT 2003 ACS on STN 118:106330 Secondary lithium batteries with coated anodes. Kanbara, Teruhisa; Sato, Yoshiko; Uemachi, Yasushi; Matsuda, Hiromu; Tonomura, Tadashi; Takeyama, Kenichi (Matsushita Electric Industrial Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 04248276 A2 **19920903** Heisei, 5 pp. (Japanese). JKXXAF. APPLICATION: JP 1991-7502 19910125. The batteries use Li+-conductive electrolytes, ABLi=-conductive cathodes, and Li (alloy) anodes coated with a substance having both Li+ and electron cond. Preferably, the coating substances are graphitic C or transition metal dichalcogenides. 12058-18-3, Molybdenum selenide (MoSe2) IT (coatings, lithium anodes contg., for secondary batteries) 12058-18-3 HCA RNMolybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNSe Mo Se

ICM H01M010-40

ICS

H01M004-02

IC

- CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
- ST lithium battery anode carbon coating; chalcogenide coating lithium anode; molybdenum sulfide coating lithium anode
- IT Transition metal chalcogenides
 (coatings, lithium anodes contg., for secondary batteries)
- IT Anodes

(battery, lithium, with lithium ion- and electron-conductive coatings, for dendrite growth prevention)

IT Lithium alloy, base

(anodes, with lithium ion- and electron-conductive coatings, for secondary batteries)

IT 7439-93-2, Lithium, uses

(anodes, with lithium ion- and electron-conductive coatings, for secondary batteries)

1317-33-5, Molybdenum sulfide (MoS2), uses 7782-42-5, Graphite, uses 12039-13-3, Titanium sulfide (TiS2) 12058-18-3, Molybdenum selenide (MoSe2) 12136-97-9, Niobium sulfide (NbS2) (coatings, lithium anodes contg., for secondary batteries)

L56 ANSWER 2 OF 18 HCA COPYRIGHT 2003 ACS on STN
114:9642 Solid-state alkali metal batteries having porous
cathode current collectors. Fauteux, Denis G. Moore,
Michael J.; Blonsky, Peter M. (USA). U.S. US 4925752 A
19900515, 7 pp. (English). CODEN: USXXAM. APPLICATION: US
1989-319434 19890303.

A laminar battery comprises an alkali metal/anode AB (Li foil, a metal foil coated with a layer of Li or Li alloy), a solid ionically conducting electrolyte layer , and a cathode/current collector layer, which consists of an elec. conductive substrate having a plurality of surface voids and a radiation-cured cathode compn. compn. comprises a compd. such as V6013, MoO2, TiS2, FeOCl; an elec. conductive filler (C particles); and a radiation-cured ionically conductive electrolyte. The electrolyte comprises a solid soln. of an ionizable alkali metal or alk. earth salt and an ionically conductive polymer contg. a repeating unit CH2C(H)(R)O, CHCH2NR1, or CH2CH(OR3R2), where R is H, R2, CH2OR2, CH2OR3R2, CH2NMe2; R1 is R2 or R3R2, R2 is C1-16 (preferably C1-4) alkyl or C5-8 cycloalkyl group and R3 is an ether group. The conductive substrate is C, Cu, Al, Ni, steel, and/or Pb. The importance of roughened surface of the conductive substrate (etched Ni foil, Ni felt) on the Li battery performance was demonstrated. The battery cathode mixt. consisted of V6013 45, C 4, and electrolyte 51%. The electrolyte consisted of propylene carbonate 70, PEO 3, LiCF3SO3 6, and radiation-curable acrylate 21%.

12034-77-4 HCA RN Niobium selenide (NbSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNSe Nb Se ICM H01M006-18 TC 429191000 NCL52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC Section cross-reference(s): 38 lithium vanadium oxide battery; cathode etched STnickel current collector; solid electrolyte lithium vanadium battery Acrylic polymers, uses and miscellaneous IT(electrolytes from lithium-PEO complexes and propylene carbonate and, for batteries) Cathodes IT (battery, vanadium oxide, contg. lithium trifluoromethanesulfonate and conductive polymer, with etched nickel current collector) 7429-90-5, Aluminum, uses and miscellaneous 7439-92-1, Lead, uses IT 7440-02-0, Nickel, uses and miscellaneous and miscellaneous 7440-44-0, Carbon, uses and miscellaneous 7440-50-8, Copper, uses and miscellaneous 12597-69-2, Steel, uses and miscellaneous (cathode collector, surface etched, for batteries) 1307-96-6, Cobalt oxide (CoO), uses and miscellaneous 1313-13-9, IT Manganese dioxide, uses and miscellaneous 1314-62-1, Vanadium 1317-37-9, Iron sulfide (FeS) oxide (V2O5), uses and miscellaneous 1317-38-0, Copper oxide (CuO), uses and miscellaneous Lithium vanadium oxide 12018-01-8, Chromium oxide (CrO2) 12033-29-3, Molybdenum sulfide (MOS3) 12034-77-4, Niobium 12039-13-3, Titanium disulfide 12137-52-9, Vanadium diselenide 12166-28-8, Vanadium disulfide 13870-10-5, Iron oxide (V3O8) 15915-20-5, Chromium oxybromide (CrOBr) chloride oxide (FeClO) 16812-54-7, Nickel sulfide (NiS) 18868-43-4, Molybdenum oxide (MoO2) (cathodes, contg. alkali metal or alk. earth salt and conductive polymer, current collectors for, in batteries 12037-42-2, Vanadium oxide (V6013) IT (cathodes, contg. lithium trifluoromethanesulfonate, current collectros for, in batteries) 108-32-7, Propylene carbonate IT(electrolytes from lithium-PEO complexes and radiation-curable acrylate and, for batteries) 25322-68-3D, PEO, lithium 7439-93-2D, Lithium, PEO complexes TTcomplexes (electrolytes from propylene carbnate and radiation-curable

acrylate and, for batteries)

```
HCA COPYRIGHT 2003 ACS on STN
    ANSWER 3 OF 18
108:207653 Studies on niobium triselenide cathode material for
     lithium rechargeable cells. Ratnakumar, B. V.; Ni, C. L.;
     DiStefano, S.; Somoano, R. B.; Bankston, C. P. (Jet Propul. Lab.,
     California Inst. Technol., Pasadena, CA, 91109, USA). Proceedings -
     Electrochemical Society, 88-6 (Proc. Symp. Primary Second. Ambient
     Temp. Lithium Batteries, 1987), 565-80 (English) 1988.
     CODEN: PESODO. ISSN: 0161-6374.
     NbSe3 cathode intercalates reversibly with 3 equivs. Li at
AΒ
     a utilization efficiency of .apprx.90\%. The Li intercalation occurs
     at different closely spaced potentials between 1500 and 1800 mV vs.
              The kinetics of NbSe3 redn. is essentially governed by the
     slow diffusion of Li ions inside the layered
     cathode. The exchange c.d. and apparent transfer coeff. are
     32 .mu.A/cm2 and 0.21, resp. The prepn., characterization, and
     performance of NbSe3 are described. Several electrochem. techniques
     (cyclic voltammetry, const. current/const. potential discharge, d.c.
     potentiodynamic scans, a.c. impedance, and a.c. voltammetry) were
     used to study the Li intercalation mechanism. A phys. model is
     presented to illustrate the various steps involved in NbSe3 redn.
     12034-78-5, Niobium triselenide
IT
        (cathodes, for lithium rechargeable batteries
     12034-78-5 HCA
RN
     Niobium selenide (NbSe3) (7CI, 9CI) (CA INDEX NAME)
CN
     Se
Se Nb Se
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
     Section cross-reference(s): 72
     niobium selenide cathode lithium battery;
ST
     intercalation cathode niobium selenide battery
     Electrolytic polarization
IT
         (of niobium selenide prismatic cathode, in lithium
        hexafluoroarsenate-methyltetrahydrofuran electrolyte)
     Cathodes
IT
         (battery, intercalation, niobium selenide, for lithium
        rechargeable batteries)
     12034-78-5, Niobium triselenide
IT
         (cathodes, for lithium rechargeable batteries
     55886-04-9P, Lithium niobium selenide (Li3NbSe3)
IT
         (formation of, in discharge of lithium-niobium triselenide
        battery cathode)
      61673-65-2P, Lithium niobium selenide
 IT
         (intercalated, formation of, in discharge of lithium-niobium
         triselenide battery cathode)
```

ANSWER 4 OF 18 HCA COPYRIGHT 2003 ACS on STN

L56

```
105:199063 Photoelectrochemical cells with n-type zinc selenide and
    n-type antimony selenide (Sb2Se3) thin film semiconductor
     electrodes. Roy, C. B.; Nandi, D. K.; Mahapatra, P. K.
     (Dep. Chem., Indian Inst. Technol., Kharagpur, India).
     Electrochimica Acta, 31(10), 1227-9 (English) 1986.
                     ISSN: 0013-4686.
     CODEN: ELCAAV.
     Photoelectrochem. cells with thin film semiconductor
AB
     electrode made of ZnSe or Sb2Se3 and a platinized Pt
     electrode as counter electrode with I-/I3-
     electrolyte were investigated. The max. efficiency was obsd. with
     the cells for light wavelength .apprx.600 nm. The conversion
     efficiencies of the cells with ZnSe electrode and Sb2Se3
     electrode were 0.03% and 0.13%, resp., with light intensity
     of 92 mW cm-2.
     1315-05-5 1315-09-9
IT
        (electrodes, photoelectrochem. semiconductive)
     1315-05-5 HCA
RN
     Antimony selenide (Sb2Se3) (8CI, 9CI) (CA INDEX NAME)
CN
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***
     1315-09-9 HCA.
     Zinc selenide (ZnSe) (9CI) (CA INDEX NAME)
CN
Se==Zn
     72-2 (Electrochemistry)
CC
     Section cross-reference(s): 52, 76
     antimony selenide photoelectrochem semiconductive electrode
ST
     ; zinc selenide photoelectrochem semiconductive electrode;
     selenide antimony zinc photoelectrochem electrode; iodide
     redox electrolyte photoelectrochem cell
     Electrodes
IT
        (photoelectrochem., semiconductive, antimony selenide and zinc
        selenide)
     7440-06-4, uses and miscellaneous
IT
        (electrode, counter, in photoelectrochem. cell with
        antimony selenide or zinc selenide semiconductive
        electrodes and iodide/triiodide redox electrolyte)
     1315-05-5 1315-09-9
IT
        (electrodes, photoelectrochem. semiconductive)
     ANSWER 5 OF 18 HCA COPYRIGHT 2003 ACS on STN
L56
            Correction of: 97:147577 Semiconductor electrodes
     in contact with aqueous and nonaqueous redox-electrolytes for
     photovoltaic solar energy conversion..
     Gerischer, H.; Decker, F.; Kautek, W. (Fritz-Haber-Inst.,
     Max-Planck-Ges., Berlin, D-1000, Fed. Rep. Ger.). Solar Energy R&D
     in the European Community, Series D: Photochemical,
     Photoelectrochemical and Photobiological Processes, 1(Photochem.,
     Photoelectrochem. Photobiol. Processes), 88-93 (English)
     1982. CODEN: SRDDD6. ISSN: 0167-7950.
```

The exceptional behavior of I-/I3- system in contact with

AΒ

transition-metal dichalcogenide (MoS2, MoSe2, and WSe2) photoelectrodes, and electrodes of semiconductor oxide heterojunctions for photoelectrolysis are discussed. SnO2- and In203-coated n-Si were tested in different aq. solns. and the chem. and electrochem. stability of such electrodes against corrosion was proved. 12058-18-3 12067-46-8 ΙT (electrodes, photoelectrochem.-cell, performance of, in ag. redox electrolytes) 12058-18-3 HCA RNMolybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNSe Mo Se 12067-46-8 HCA RNTungsten selenide (WSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CN Se == W == Se 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CCSection cross-reference(s): 76 photoelectrochem cell electrode molybdenum sulfide; ST selenide molybdenum photoelectrochem cell; tungsten selenide photoelectrochem cell; silicon tin oxide photoelectrochem cell; indium oxide silicon photoelectrochem cell; iodide photoelectrochem cell electrolyte IT Electrodes (photoelectrochem., semiconductive, performance of, in aq. and nonaq. redox electrolytes) 7440-21-3, uses and miscellaneous IT (electrodes from indium oxide- or tin oxidecoated, photoelectrochem.-cell, performance of) 18282-10-5 1312-43-2 IT (electrodes from silicon coated with, photoelectrochem.-cell, performance of) 12058-18-3 12067-46-8 IT(electrodes, photoelectrochem.-cell, performance of, in aq. redox electrolytes) 14900-04-0 IT(electrolyte contg., photoelectrochem.-cell, performance of molybdenum and tungsten chalcogenide electrodes in aq.) ANSWER 6 OF 18 HCA COPYRIGHT 2003 ACS on STN 104:92052 Optical energy conversion in photoelectrochemical cells with semiconductor electrodes. Maruyama, Toshiro; Goto, Kazuhito (Coll. Eng., Kyoto Univ., Kyoto, Japan). Kagaku Kogaku, 50(1), 58-61 (Japanese) 1986.

ISSN: 0375-9253.

Polycryst. semiconductor films for regeneration-type

CODEN: KKGKA4.

AB

photoelectrochem. cells were made of CdSe and TiO2 by electrophoretic pptn. The current-potential property and solar energy conversion efficiency of the thin films as anodes were studied.

RN 1306-24-7 HCA

CN Cadmium selenide (CdSe) (9CI) (CA INDEX NAME)

Cd=== Se

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

ST cadmium selenide photoelectrochem **electrode**; titania photoelectrochem **electrode**

IT Anodes

(photoelectrochem., cadmium selenide and titania, prepn. and properties of)

IT 1306-24-7P, uses and miscellaneous 13463-67-7P, uses and miscellaneous

(electrodes, photoelectrochem., prepn. and properties of)

L56 ANSWER 7 OF 18 HCA COPYRIGHT 2003 ACS on STN

101:100115 Semiconductor-electrolyte photoelectrode systems. IV.
Electrodeposition and photoelectrochemical properties of manganese selenide (MnSe) film electrode. Song, Tianping;
Wang, Yujiang; Lu, Wenzhe; Yang, Jun; Mi, Tianying (Changchun Inst. Appl. Chem., Acad. Sin., Changchun, Peop. Rep. China). Zhongguo Kexueyuan Changchun Yingyong Huaxue Yanjiuso Jikan, 19, 34-42 (Chinese) 1982. CODEN: ZKCJEE.

The bandgap of the electrodeposited MnSe was estd. by photoacoustic measurements to be about 1.91 eV in agreement with that reported in literature. Photoelectrochem. measurements showed typical photovoltaic characteristics with energy conversion efficiency about 1% under Xe light of moderate

conversion efficiency about 1% under Xe light of moderate
intensity. Doping (by codeposition) of certain rare-earth (such as
Pr, Sm, etc.) ions considerably enhanced the photoeffects of the
MnSe electrode.

IT 37320-90-4

(electrodes, photoelectrochem.)

RN 37320-90-4 HCA

CN Manganese selenide (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
Se Mn	x x x	7782-49-2 7439-96-5

CC 72-2 (Electrochemistry)

Section cross-reference(s): 76 manganese selenide photoelectrochem electrode; rare earth STdoping manganese selenide Rare earth metals, uses and miscellaneous IT(doping with, of manganese selenide photoelectrochem. electrodes) Electrodes IT (photoelectrochem., manganese selenide) 37320-90-4 IT (electrodes, photoelectrochem.) ANSWER 8 OF 18 HCA COPYRIGHT 2003 ACS on STN L56 99:215705 Electrochemical solar cells with layer-type semiconductor anodes. Stabilization of the semiconductor electrode by selective polyindole electrodeposition. Fornarini, L.; Stirpe, F.; Scrosati, B. (Ist. Chim. Fis., Univ. Rome, Rome, Italy). Journal of the Electrochemical Society, 130(11), 2184-7 (English) 1983. ISSN: 0013-4651. CODEN: JESOAN. Electropolymn. of indole was investigated on Pt and MoSe2 ABelectrodes. When performed in the dark, the electropolymn. is selectively directed to the surface defects of the semiconductor. The effect of this surface treatment was investigated. The electropolymn. of indole effectively blocks the defect sites and produces a stable improvement on the output characteristics of photoelectrochem. cells with layer-type semiconductor anodes. 12058-18-3 IT(anodes from polyindole-stabilized, photoelectrochem.-cell) 12058-18-3 HCA RN Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CNSe Mo Se 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CCSection cross-reference(s): 38, 72 platinum polyindole electrode; molybdenum selenide ST polyindole photoelectrochem cell; electrode molybdenum selenide polyindole photoelectrochem IT Anodes (photoelectrochem., molybdenum selenide, polyindole-stabilized) 12058-18-3 IT (anodes from polyindole-stabilized, photoelectrochem.-cell) IT 82451-55-6 (anodes stabilized with, layer-type semiconductor, photoelectrochem.-cell) 7440-06-4, uses and miscellaneous IT (electrodes, contg. deposited polyindole, voltammogram of, in aq. iodide-iodine)

```
ANSWER 9 OF 18 HCA COPYRIGHT 2003 ACS on STN
L56
98:92635 Electrochemical solar cells using cadmium
     selenide thin-film electrodes. Xiao, Xu Rui;
     Tien, H. Ti (Dep. Biophys., Michigan State Univ., East Lansing, MI,
     48824, USA). Journal of the Electrochemical Society, 130(1), 55-9
     (English) 1983. CODEN: JESOAN. ISSN: 0013-4651.
     Electrochem. photocells of a CdSe thin-film anode
AΒ
     and a Pt cathode immersed in M Na2S-NaOH-S soln. were
     studied. CdSe thin films were formed on Ti, Cr, Mo, SnO2,
     glassy C, and graphite substrates by coating an aq. mixt.
     of CdSe, ZnCl2, and a surfactant and subsequently sintering at
     400-500.degree. in air. The current-voltage (I-V) relations, output
     power efficiency, open-circuit voltage, and short-circuit current
     were measured. A 7% power conversion efficiency was obtained at 20
     mW/cm2 light intensity after photoetching. The monochromatic I-V
     curves were analyzed.
     1306-24-7, uses and miscellaneous
IT
        (anodes, photoelectrochem.-cell, performance of thin-
        film)
     1306-24-7 HCA
RN
     Cadmium selenide (CdSe) (9CI) (CA INDEX NAME)
CN
Cd== Se
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
     photoelectrochem cell cadmium selenide electrode; titanium
ST
     cadmium selenide photoelectrochem electrode; chromium
     cadmium selenide photoelectrochem electrode; molybdenum
     cadmium selenide photoelectrochem electrode; carbon
     cadmium selenide photoelectrochem electrode; graphite
     cadmium selenide photoelectrochem electrode; tin dioxide
     cadmium selenide electrode
     Anodes
IT
     Photoelectric devices, solar
        (photoelectrochem., cadmium selenide, performance of thin-
     1306-24-7, uses and miscellaneous
ΙT
        (anodes, photoelectrochem.-cell, performance of thin-
     7439-98-7, uses and miscellaneous 7440-32-6, uses and
TT
                    7440-47-3, uses and miscellaneous
                                                        7782-42-5, uses
     miscellaneous
                         18282-10-5
     and miscellaneous
        (electrodes from cadmium selenide-coated,
        photoelectrochem.-cell, performance of thin-film)
     7440-44-0, uses and miscellaneous
ΙT
         (glassy, electrodes from cadmium selenide-
        coated, photoelectrochem.-cell, performance of thin-
        film)
```

ANSWER 10 OF 18 HCA COPYRIGHT 2003 ACS on STN

```
97:147577 Semiconductor electrodes in contact with aqueous and
     nonaqueous redox-electrolytes for photovoltaic solar energy
                 Gerischer, H.; Decker, F.; Kautek, W.
     conversion.
     (Fritz-Haber-Inst., Max-Planck-Ges., Berlin, D-1000/33, Fed. Rep.
             Solar Energy R&D in the European Community, Series D:
     Photochemical, Photoelectrochemical and Photobiological Processes,
     1(Photochem., Photoelectrochem. Photobiol. Processes), 88-93
     (English) 1982. CODEN: SRDDD6. ISSN: 0167-7950.
     The exceptional behavior of I-/I3- system in contact with
AΒ
     transition-metal dichalcogenide (MoS2, MoSe2, and WSe2)
     photoelectrodes, and electrodes of semiconductor oxide
     heterojunctions for photoelectrolysis are discussed. SnO2- and
     Tn203-coated n-Si were tested in different aq. solns. and
     the chem. and electrochem. stability of such electrodes
     against corrosion was proved.
     12058-18-3 12067-46-8
IT
        (electrodes, photoelectrochem.-cell, performance of, in
        aq. redox electrolytes)
     12058-18-3 HCA
RN
     Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
Se Mo Se
     12067-46-8 HCA
RN
     Tungsten selenide (WSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
Se = W = Se
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
     Section cross-reference(s): 76
     photoelectrochem cell electrode molybdenum sulfide;
ST
     selenide molybdenum photoelectrochem cell; tungsten selenide
     photoelectrochem cell; silicon tin oxide photoelectrochem cell;
     indium oxide silicon photoelectrochem cell; iodide photoelectrochem
     cell electrolyte
     Electrodes
ΙT
        (photoelectrochem., semiconductive, performance of, in aq. and
        nonaq. redox electrolytes)
     7440-21-3, uses and miscellaneous
IT
        (electrodes from indium oxide- or tin oxide-
        coated, photoelectrochem.-cell, performance of)
     1312-43-2
                 18282-10-5
IT
        (electrodes from silicon coated with,
        photoelectrochem.-cell, performance of)
     12058-18-3 12067-46-8
ΙT
        (electrodes, photoelectrochem.-cell, performance of, in
        aq. redox electrolytes)
ΙT
     14900-04-0
        (electrolyte contg., photoelectrochem.-cell,
        performance of molybdenum and tungsten chalcogenide
```

electrodes in aq.)

L56 ANSWER 11 OF 18 HCA COPYRIGHT 2003 ACS on STN 96:202533 Electrochemical solar cells with layer-type semiconductor anodes. Nonaqueous electrolyte cells. Fornarini, L.; Stirpe, F.; Scrosati, B. (Ist. Chim. Fis., Univ. Rome, Rome, Italy). Journal of the Electrochemical Society, 129(5), 1155-6 (English) 1982. CODEN: JESOAN. ISSN: 0013-4651.

The output characteristics of the n-MoSe2 3 KI, 0.15M I2, formamide Pt cell are promising, showing fill-factor and efficiency values which approach those obtained in aq. solns. No degrdn. in the cell was obsd. after continuous operation at the max. power point for >1 wk.

IT 12058-18-3

(electrodes, photoelectrochem. cells contg., performance of nonaq.)

RN 12058-18-3 HCA

CN Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

Se = Mo = Se

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology)

IT 12058-18-3

(electrodes, photoelectrochem. cells contg., performance of nonaq.)

L56 ANSWER 12 OF 18 HCA COPYRIGHT 2003 ACS on STN

96:151221 Semiconductor electrodes. 44.

Photoelectrochemistry at polycrystalline p-type tungsten selenide
films. Abruna, Hector D.; Bard, Allen J. (Dep. Chem., Univ.

Texas, Austin, TX, 78712, USA). Journal of the Electrochemical Society, 129(3), 673-6 (English) 1982. CODEN: JESOAN.

ISSN: 0013-4651.

The title **electrodes** were used in the study of photoelectrochem. redox behavior of several common couples. The highest output was obtained with FeL3+ where L is macrocyclic ligand prepd. by condensation of diacetylpyridine and triethylenetetramine. The p-WSe2 **electrode** was characterized by the flatband potential, the doping d., the **film** resistivity and the scanning electron microscope. The use of some of these systems in solar **energy conversion** was considered.

IT 12067-46-8

(films of polycryst. p-type, photoelectrochem. of)

RN 12067-46-8 HCA

CN Tungsten selenide (WSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

Se== W== Se

CC 72-2 (Electrochemistry)

139-13-9

1824-81-3

Section cross-reference(s): 52, 76 tungsten selenide polycrystn semiconductor electrode; STphotoelectrochem redox tungsten selenide; solar energy conversion tungsten selenide ITPhotoelectric emission (from tungsten selenide films, polycryst. p-type) IT(photoelectrochem., film, tungsten selenide, polycryst. p-type) 12067-46-8 ΙT (films of polycryst. p-type, photoelectrochem. of) ANSWER 13 OF 18 HCA COPYRIGHT 2003 ACS on STN 96:71834 Electrochemical solar cells with layer-type semiconductor anodes: chemical treatments of the crystal surface. Razzini, G.; Bicelli, L. Peraldo; Pini, G.; Scrosati, B. (Ist. Elettrochim. Chim.-Fis. Metall., Politech. Milan, Milan, Italy). Journal of the Electrochemical Society, 128(10), 2134-7 (English) 1981. ISSN: 0013-4651. CODEN: JESOAN. Photoelectrochem. cells based on layer-type semiconductors AB (i.e., transition-metal sulfides and selenides) have gained substantial interest due to their reasonably high solar efficiency and good stability against photocorrosion. However, the performance of this cell type may be limited by irregularities in the semiconductor, since edges of the van der Waals surface exposed to the electrolyte may act as recombination centers. To control this effect, chem. surface treatments, based on specific agents which perform selectively on the transition metal atoms at the edge sites, were studied. In particular, the effect of the disodium salt of EDTA (as well as that of many other org. mols.) on the performance of an n-MoSe2/I-, I2/Pt sample cell was investigated. Treatment with EDTA generally improved both short-circuit current and power output, even if a considerable variation in the response from crystal to crystal was obsd. The stability of the EDTA effect under prolonged cell operation was also examd. 12058-18-3 IT (electrodes, photoelectrochem. cells contg. chem. surface-treated, performance of) 12058-18-3 HCA RNMolybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME) CN Se Mo Se 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC IT12058-18-3 (electrodes, photoelectrochem. cells contg. chem. surface-treated, performance of) 100-69-6 110-86-1, uses and miscellaneous 462-08-8 482-54-2 504-29-0 1122-58-3

IT

100-43-6

139-33-3

25014-15-7

25232-41-1

(photoelectrochem.-cell **electrodes** from molybdenum diselenide surface treated with, performance of)

L56 ANSWER 14 OF 18 HCA COPYRIGHT 2003 ACS on STN 95:190018 Electrochemical solar cells with layer-type semiconductor anodes. Performance of n-molybdenum selenide (MoSe2) cells. Razzini, G.; Lazzari, M.; Bicelli, L. Peraldo; Levy, F.; De Angelis, L.; Galluzzi, F.; Scafe, E.; Fornarini, L.; Scrosati, B. (Cent. Studio Processi Elettrodici, Polytech. Milan, Milan, Italy). Journal of Power Sources, 6(4), 371-82 (English) 1981. CODEN: JPSODZ. ISSN: 0378-7753.

The output characteristics and the long-term performances of n-MoSe2 (I-,I2) electrochem. solar cells were investigated. The surface state of the semiconductor plays a key role in the behavior of the cell. With smooth crystal samples, fill factor and efficiency values of 0.6 and 6%, resp., were obtained under air-mass-1 illumination. Such performances are, however, drastically reduced if irregular crystal samples are used. Control of these undesirable surface-state effects was attempted by chem. treatments specific to the unsatd. transition-metal atoms exposed to the electrolyte at the edge sites. The stability of n-MoSe2 (I-,I2) cells under long-time operation, was also evaluated.

IT 12058-18-3

(anodes, photoelectrochem. cells contg., long-term performance of)

RN 12058-18-3 HCA

CN Molybdenum selenide (MoSe2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

Se = Mo = Se

L56 ANSWER 15 OF 18 HCA COPYRIGHT 2003 ACS on STN 95:177608 Thin film cadmium selenide electrodes for backwall photoelectrochemical cells. Russak, Michael A.; Reichman, Joseph (Res. Dep., Grumman Aerosp. Corp., Bethpage, NY, 11714, USA). Journal of the Electrochemical Society, 128(9), 2029-31 (English) 1981. CODEN: JESOAN. ISSN: 0013-4651.

These CdSe electrodes were deposited on SnO2coated glass and have resistivity in the range of 10
.OMEGA./square and visible transmission of .apprx.80%. These
films were deposited by simultaneous vacuum evapn. of Cd and
Se onto chem.-spray-deposited, halogen-doped, SnO2-coated
glass slides. The irradn. of the CdSe was through the substrate
glass and the transparent conductor. This cell eliminates
the electrolyte absorption loss and allows reasonably
quick sample change. Current-voltage curves are given and with a
Se/Cd ratio of 3 with heat treatment in air at 400.degree. these

backwall electrodes gave power conversion efficiencies IT 79497-64-6 (electrodes, photoelectrochem.) 79497-64-6 HCA RNCadmium selenide (CdSe3) (9CI) (CA INDEX NAME) Se-Se Se-Cd 1306-24-7, uses and miscellaneous IT (electrodes, thin-film, for backwall photoelectrochem. cells) 1306-24-7 HCA RNCadmium selenide (CdSe) (9CI) (CA INDEX NAME) CNCd Se CC72-7 (Electrochemistry) Section cross-reference(s): 74, 76 cadmium selenide electrode stannic oxide; photoelectrochem STelectrode cadmium selenium; energy conversion solar elec ITElectrodes (photoelectrochem., film, cadmium selenide, for backwall cells) 79497-64-6 IT (electrodes, photoelectrochem.) 1306-24-7, uses and miscellaneous IT (electrodes, thin-film, for backwall photoelectrochem. cells) ANSWER 16 OF 18 HCA COPYRIGHT 2003 ACS on STN 95:177607 Properties of cadmium selenide thin film electrodes for photoelectrochemical cells. Reichman, Joseph; Russak, Michael A. (Res. Dep., Grumman Aerosp. Corp., Bethpage, NY, 11714, USA). Journal of the Electrochemical Society, 128(9), 2025-9 (English) 1981. CODEN: JESOAN. 0013-4651. Recent efforts on establishing the relations among thin-film ABprocessing parameters, electronic properties, and photoelectrochem. performance of CdSe films are summarized. The electrodes were deposited onto Ti sheet and were designed for use in frontwall illuminated electrochem. photovoltaic cells where the incoming radiation would pass through a transparent window, a thin layer of electrolyte, and then strike the semiconductor electrolyte junction. The most promising results were obtained with CdSe films deposited at

.apprx.100.degree. with a Se/Cd ratio of 3 and then heat-treated in the temp. range of 350-400.degree. in air. 79497-64-6 ΙT (electrodes, photoelectrochem.) RN 79497-64-6 HCA Cadmium selenide (CdSe3) (9CI) (CA INDEX NAME) CNSe-Se Se--- Cd 1306-24-7, uses and miscellaneous IT (electrodes, thin-film, for photoelectrochem. cells) 1306-24-7 HCA RN Cadmium selenide (CdSe) (9CI) (CA INDEX NAME) CNCd Se 72-7 (Electrochemistry) CC Section cross-reference(s): 74, 76 cadmium selenide film electrode STphotoelectrochem; charge transfer cadmium selenide electrode Electron exchange IT(on cadmium selenide thin-film electrodes, in photoelectrochem. cells) IT Electrodes (photoelectrochem., film, cadmium selenide) 79497-64-6 IT (electrodes, photoelectrochem.) 1306-24-7, uses and miscellaneous IT(electrodes, thin-film, for photoelectrochem. cells) ANSWER 17 OF 18 HCA COPYRIGHT 2003 ACS on STN 95:153806 Electrochemical photovoltaic cells cadmium selenide thin film electrodes. Russak, M. A.; Reichman, J.; DeCarlo, J.; Creter, C. (Res. Dep., Grumman Aerosp. Corp., Bethpage, NY, USA). Report, SERI-TR-8002-8-T1, 63 pp. Avail. NTIS From: Energy Res. Abstr. 1981, 6(11), Abstr. No. 15465 (English) 1980. Progress on developing stable, thin-film CdSe AΒ electrodes with sunlight conversion efficiency of 10% for use with aq. polysulfide electrolytes in frontwall and backwall illuminated electrochem. photovoltaic cells is reported. The relation among thin-film processing, resultant electronic properties, and current-voltage (I-V) performance was studied to produce electrodes with max. power conversion efficiency. The best results were obtained with

CdSe thin-film electrodes produced in 2 ways for frontwall cells. Films were deposited on Ti at .apprx.100.degree. with a high Se/Cd ratio and then heat treated in air at 350-400.degree.. These films usually have a very fine grained microstructure after heat treatment and the resultant electrodes exhibit fairly square I-V characteristics, with fill factors of .gtoreq.0.6 and high current output. The overall power efficiency of these electrodes is limited by relatively low output voltages. At present, power conversion efficiencies of 3-5% can be obtained reproducibly at simulated air-mass-2 conditions with electrodes processed in this The 2nd type of film that yielded promising results was deposited on Ti at substrate temp. of >400.degree.. These electrodes have increased open-circuit voltage. However, the current output and fill factor are lower. As a result, the power conversion efficiency of these electrodes is 3 to 4%. Backwall electrodes with an efficiency of >4% and short-circuit densities near theor. for air-mass-2 conditions were produced. 1306-24-7, uses and miscellaneous (electrodes, photoelectrochem. cells contg. aq.

IT

polysulfide electrolyte and, development and properties of)

RN1306-24-7 HCA

Cadmium selenide (CdSe) (9CI) (CA INDEX NAME) CN

Cd==Se

52-2 (Electrochemical, Radiational, and Thermal Energy Technology) CC ST cadmium selenide electrode photoelectrochem cell

IT Electrodes

(photoelectrochem., cadmium selenide, development and properties of)

1306-24-7, uses and miscellaneous IT

(electrodes, photoelectrochem. cells contg. aq. polysulfide electrolyte and, development and properties of)

ANSWER 18 OF 18 HCA COPYRIGHT 2003 ACS on STN

92:96735 Electrochemical photovoltaic cells

utilizing thin film semiconductor electrodes.

Russak, M. A.; Reichman, J.; Witzke, H.; Deb, S. K.; Chen, S. N. (Res. Dep., Grumman Aerosp. Corp., Bethpage, NY, 11714, USA). Comm. Eur. Communities, [Rep.] EUR, EUR 6376, Photovoltaic Sol. Energy Conf., 690-700 (English) 1979. CODEN: CECED9.

The fabrication and evaluation of thin-film CdSe AΒ electrodes for use in the title cells is described. effect of electrode prepn. and electrolyte concn. on efficiency and stability are presented. Current-voltage curves and action spectra are also discussed, and a relation between efficiency and minority carrier diffusion length is shown. Conversion efficiencies as high as 5% are reported.

1306-24-7P, uses and miscellaneous ΙT

```
(electrodes, photoelectrochem.-cell, prepn. and
        properties of thin-film)
     1306-24-7 HCA
RN
     Cadmium selenide (CdSe) (9CI) (CA INDEX NAME)
CN
Cd== Se
     52-2 (Electrochemical, Radiational, and Thermal Energy Technology)
CC
     photoelectrochem solar cell electrode; selenide cadmium
ST
     electrode photoelectrochem cell
     Electrodes
IT
        (photoelectrochem.-cell, cadmium selenide, prepn. and properties
        of thin-film)
     Electric current carriers
IT
        (minority, diffusion length of, efficiency of cadmium selenide
        photoelectrochem. cell electrodes in relation to)
IT
     1306-24-7P, uses and miscellaneous
        (electrodes, photoelectrochem.-cell, prepn. and
        properties of thin-film)
=> d 157 1-8 cbib abs hitstr hitind
     ANSWER 1 OF 8 HCA COPYRIGHT 2003 ACS on STN
127:284789 Thermal spray deposited electrode component and
     method of manufacture. Muffoletto, Barry C.; Paulot, William M.;
     Spaulding, Joseph E. (Wilson Greatbatch Ltd., USA). PCT Int. Appl.
     WO 9736023 A1 19971002, 42 pp. DESIGNATED STATES: W: AU,
     JP; RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL,
              (English). CODEN: PIXXD2. APPLICATION: WO 1996-US17558
     19961030. PRIORITY: US 1996-621257 19960325.
     An electrode component for an electrochem.
AB
     cell is described wherein the electrode is
     produced by thermal spraying an electrode active material
     onto a substrate to coat the substrate. Suitable thermal spraying
     processes include chem. combustion spraying and elec. heating
     spraying, using both wire and power processes.
     12039-13-3, Titanium disulfide
IT
        (cathode active material; electrochem.
        cell with cathode coated with)
RN
     12039-13-3 HCA
     Titanium sulfide (TiS2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
s = Ti = s
TC:
     I.CM
          C25B009-00
          C25B011-04; C25B011-10; C23C004-10; H01M006-04; H01M006-14;
     ICS
          H01M006-00; B23P019-00
     72-2 (Electrochemistry)
CC
     thermal spray deposited electrode component;
ST
```

electrolytic cell electrode Alkali metals, uses IT(electrochem. cell with anode from) Electrolytic cells IT(electrode for) Electrodes IT(thermal spray deposited electrode component and method of manuf.) Coating process IT (thermal spraying; thermal spray deposited electrode component and method of manuf.) 1313-99-1, Nickel oxide, uses 1313-13-9, Manganese dioxide, uses IT7440-44-0, Carbon, uses 7440-44-0D, 1344-70-3, Copper oxide Carbon, fluorinated, uses 11104-61-3, Cobalt oxide 11115-78-9, 11118-57-3, Chromium oxide 11126-12-8, Iron Copper sulfide sulfide 12039-13-3, Titanium disulfide 181183-66-4, Copper silver vanadium oxide (cathode active material; electrochem. cell with cathode coated with) 7439-93-2, Lithium, uses IT(electrochem. cell with anode from) 16919-18-9D, Hexafluorophosphate, alkali metal salt 16973-45-8D, ITHexafluoroarsenate, alkali metal salt (electrochem. cell with electrolyte from) 67-68-5, Dimethyl sulfoxide, uses 68-12-2, Dimethyl formamide, uses 75-05-8, Acetonitrile, uses 79-20-9, Methyl acetate IT96-49-1, Ethylene carbonate 109-99-9, Tetrahydrofuran, uses 112-49-2, Triglyme 127-19-5, Dimethyl 111-96-6, Diglyme acetamide (electrochem. cell with electrolyte in solvent from) 108-32-7, Propylene carbonate 110-71-4 IT (electrochem. cell with electrolyte in solvent from mixt. of dimethoxyethane and propylene carbonate) 7440-32-6, Titanium, uses 7429-90-5, Aluminum, uses IT(electrode substrate; silver vanadium oxide-coated electrode substrate for electrochem. cell) 11105-02-5, Silver vanadium oxide IT (silver vanadium oxide-coated electrode for

ANSWER 2 OF 8 HCA COPYRIGHT 2003 ACS on STN 127:114512 Durable electrode coatings. Tsou, Yu-min (Dow

electrochem. cell)

Chemical Co., USA). U.S. US 5645930 A 19970708, 15 pp. (English). CODEN: USXXAM. APPLICATION: US 1995-513581 19950811.

Durable electrolytic cell electrodes AΒ having low hydrogen overpotential and performance stability are presented. A highly porous electrocatalytic primary phase and an outer, secondary phase reinforcement coating are provided on an elec. conducting transition metal substrate to make the

electrodes. Durability is achieved by the application of the outer secondary phase to protect the primary phase electrocatalytically active coating. A process is also disclosed for catalyzing a substrate surface to promote electroless deposition of a metal.

IT 11113-75-0, Nickel sulfide

(plating of nickel sulfide on polycarbonate in manufg. durable electrode)

RN 11113-75-0 HCA

CN Nickel sulfide (9CI) (CA INDEX NAME)

	mponent	Ratio	Component Registry Number
S	=========	x	7704-34-9
Ni		X	7440-02-0
IC	ICM B32B0		
NCL	428328000		
CC	72-2 (Electrochemistry)		
ST	electrolytic cell electrodes low		
*		overpotential	
IT	Coating ma		
		le electrode coatings	<u> </u>
IT	Brines (durable electrode coatings for electrolysis of)		
			s for electrolysis of
IT	Overvolta	ge	with less hardwaren
	(durab	le electrode coatings	with low hydrogen
T 00		tential and performar	ice stability)
IT	Electrode	s rolytic cell electrod	log
	having	low hydrogen overnot	tential and performance stability)
ΙT		nates, uses	telletat and performance beautiful,
Т.1	forycarbo.	rate: plating of nick	cel phosphide on polycarbonate in
	manufa	. durable electrode)	· ·
IT		, Copper, uses	
	(durab	le electrode coatings	s prepd. by dipping copper
		creen in coating solr	
IT	7439-89-6	. Iron, uses	
	(durab	le electrode coatings	s prepd. by dipping iron wire
•		in coating solns.)	,
IT	11129-89-	8, Platinum oxide	1 1 1' '
	(durab	le electrode coatings	s prepd. by dipping metal
	wire s	creen in soln. contg	.)
\cdot IT	7647-01-0	, Hydrochloric acid,	uses 7647-10-1, Palladium dichloride uses 9060-90-6, Poly(aminostyrene)
	7718-54-9	, Nickel dichioride,	ride 10139-58-9, Rhodium nitrate
	10049-08-	8, Ruthenium trichlo	
	11113-77-	2, Palladium oxide	12036-10-1, Ruthenium dioxide
	12645-46-	4, Iridium oxide I.	2648-47-4, Platinum chloride 5014-15-7, Poly(2-vinylpyridine)
	12680-36-	3, Rhodium oxide 2	DOTA-TO-1' LOTA (7 ATHATATIO)

25014-41-9, Poly(acrylonitrile) 25067-59-8, Poly(vinylcarbazole)

Maples 09/432,334 25067-61-2, Poly(methacrylonitrile) 25232-41-1, 30551-89-4, Poly(allylamine) 55917-50-5, Poly(4-vinylpyridine) 61970-39-6, Osmium oxide Palladium phosphate (durable electrode coatings prepd. by dipping nickel wire screen in soln. contg.) 7440-22-4, Silver, uses (durable electrode coatings prepd. by dipping silver wire screen in coating solns.) 12597-68-1, Stainless steel, uses (durable electrode coatings prepd. by dipping stainless steel wire screen in coating solns.) 1333-74-0, Hydrogen, properties (durable electrode coatings with low hydrogen overpotential and performance stability) 7440-02-0, Nickel, uses (electrode; durable electrode coatings prepd. by dipping nickel wire screen in coating solns.) 12619-68-0, Cobalt boride (plating of Co boride on polycarbonate in manufg. durable electrode) 7440-48-4, Cobalt, properties (plating of Co on polycarbonate in manufg. durable electrode) 12643-12-8, Cobalt phosphide (plating of Co/P alloy on polycarbonate in manufg. durable electrode) 12619-90-8, Nickel boride (plating of nickel boride on polycarbonate in manufg. durable electrode)

IT

11104-08-8, Nickel phosphide IT(plating of nickel phosphide on polycarbonate in manufg. durable electrode)

11113-75-0, Nickel sulfide IT(plating of nickel sulfide on polycarbonate in manufg. durable electrode)

ANSWER 3 OF 8 HCA COPYRIGHT 2003 ACS on STN L57 106:199242 Cathodes. Kanebori, Keiichi; Kirino, Fumiyoshi; Hiratani, Masahiko; Ito, Yukio; Miyauchi, Katsumi; Kudo, Tetsuichi (Hitachi, Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 61256563 A2 19861114 Showa, 4 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1985-97761 19850510.

Conductive and heat-resistant macromol. coatings are formed between ABthe substrate and metal chalcogenide active material in battery cathodes. Polyimide resin soln. contg. dispersions of Ni, Ti, W, and graphite was spin-coated on a stainless steel substrate and heated at 200 and 350.degree. to give an .apprx.10-.mu. coating. Thin TiS2 film was formed on the coating by plasma chem. vapor deposition from TiCl4 and H2S. The resistance between the substrate and TiS2 was .ltoreq.100.OMEGA. vs. .gtoreq.10 M.OMEGA. for cathode without the polyimide interlayer.

12039-13-3, Titanium disulfide IT

ΙT

IT

IT

IT

IT

IT

IT

(cathodes with stainless steel grids coated with metal-contg. polyimides, for batteries)

RN 12039-13-3 HCA

CN Titanium sulfide (TiS2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

s = Ti = s

IC ICM H01M004-02

CC 52-2 (Electrochemical, Radiational, and Thermal Energy Technology) Section cross-reference(s): 35, 38

cathode battery polyimide metal coating; sulfide titanium cathode polyimide; nickel polyimide coating battery cathode; titanium polyimide coating battery cathode; tungsten polyimide coating battery cathode; graphite polyimide coating battery cathode; elec resistance polyimide coating cathode

IT Coating materials

(polyimides, contg. metals, for stainless steel grids, for titanium sulfide battery cathodes)

IT Cathodes

(battery, titanium sulfide, with stainless steel grids coated with metal-contg. polyimides)

IT 12039-13-3, Titanium disulfide

(cathodes with stainless steel grids coated with metal-contg. polyimides, for batteries)

IT 12597-68-1

(coating materials, polyimides, contg. metals, for stainless steel grids, for titanium sulfide **battery** cathodes)

TT 7440-02-0, Nickel, uses and miscellaneous 7440-32-6, Titanium, uses and miscellaneous 7440-33-7, Tungsten, uses and miscellaneous 7782-42-5, Graphite, uses and miscellaneous (in polyimide films, on stainless steel grids, for titanium sulfide battery cathodes)

L57 ANSWER 4 OF 8 HCA COPYRIGHT 2003 ACS on STN 91:184061 Cathode for brine electrolysis. Kajiyama, Yoshihisa; Murakami, Yoshio; Matsuura, Shunji (Tokuyama Soda Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 54087680 19790712 Showa, 3 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1977-155628 19771226.

In prepg. a cathode for an electrolysis cell by sinter coating an Fe base material with a cathode-active material, the Fe base material is pretreated in a soln. contg. .gtoreq.1 phosphates selected from Mn phosphates, Zn phosphates, or Fe phosphates. The cathode has a lower H overvoltage compared to a conventionally prepd. electrode without this pretreatment. Thus, a 10 x 30 mm steel sheet (5541) was polished with emery paper, rinsed, dipped in 10% HCl, dipped for 10 min in a 60.degree. bath contg. H3PO4 60, Zn3(PO4)2.4H2O 10,

هن 🕴 (ب

NaH2PO4.2H2O 10 g, and H2O 3%, coated with a suspension contg. Na(SCN)2 40, Me cellulose 1.5, poly(ethylene glycol) 1.5, and H2O 30 parts, then heated for 1 h at 900.degree. in an inert atm., the final coating and heating operation being repeated 4 times. ΙT 12030-51-2 16812-54-7 (in coating, sinter, of phosphate treated steel cathodes 12030-51-2 HCA RNIridium sulfide (IrS2) (7CI, 9CI) (CA INDEX NAME) CNs = Ir = S16812-54-7 HCA RN Nickel sulfide (NiS) (8CI, 9CI) (CA INDEX NAME) CNNi = SICC25B011-04 72-10 (Electrochemistry) CC cathode steel phosphating sinter coating; nickel sinter STcoating steel cathode ΙT Brines (electrolysis of, cathodes for) Cathodes IT (steel, phosphating and sinter coating of) Coating process IT(sintering, of phosphated steel cathodes) 12597-69-2, uses and miscellaneous ΙT (cathodes, phosphating and sinter coating of) 10026-00-3 **12030-51-2** 13465-52-6 6010-09-9 IT13689-92-4 **16812-54-7** 57229-17-1 (in coating, sinter, of phosphate treated steel cathodes) L57 ANSWER 5 OF 8 HCA COPYRIGHT 2003 ACS on STN 80:33330 Electrode. Chisholm, Raymond S. (Nora International Co.). U.S. US 3770613 19731106, 6 pp. Division of U.S. 3,649,485 (CA 76;135112c). (English). CODEN: USXXAM. APPLICATION: US 1971-104454 19710106. The electrodes have a coating applied to an AΒ electroconductive base. The coating consists of the sulfides, nitrides, borides and carbides of Al, Ta, Ti, Bi, W, Zr, and Hf mixed with the metals, oxides, sulfides, nitrides, borides and carbides of Au, Ag, Pt, Pd, Ru, Rh, Ir, Os, Ni, Cr, Pb, Cu and Mn. The use of the novel electrodes in alkali metal-Cl cells, both diaphragm and Hg type, alkali metal chlorate cells and other similar electrolytic applications is discussed. A coating compn. is prepd. by mixing toluene soln. of Pt resinate (7.5 wt. % Pt) 3.75, Ti sulfide 1, and toluene 4 g. The Ti sulfide is thoroughly mixed in the toluene-resinate mixt., and the resulting mixt. is painted on a Ti strip which is, prior to painting, pickled in HCl soln. The painted surface is heated in air to 450.degree. for 1 hr. The procedure is repeated 5 times to provide a tightly bonded coating of Ti sulfide-Pt to the Ti base.

IT 12038-21-0 12673-92-6

(coating with, of graphite and titanium electrodes, for brine electrolysis)

RN 12038-21-0 HCA

CN Platinum sulfide (PtS2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)

S = Pt = S

RN 12673-92-6 HCA

CN Titanium sulfide (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
======================================	+=====================================	7704-34-9 7440-32-6

IC B01K

NCL 204290000R

CC 77-10 (Electrochemistry)

ST electrode electrolytic cell chlorine chlorate; alkali metal chlorate cell electrode; coating titanium base electrode

IT Brines

(electrolysis of, coating of graphite and titanium electrodes for)

IT Electrodes

(for brine electrolysis, coating of graphite and titanium)

IT Chlorates

(manuf. of, coating of graphite and titanium electrodes for)

IT Coating process

(of graphite and titanium **electrodes**, for brine electrolysis)

IT 11113-84-1 **12038-21-0** 12070-08-5 12673-91-5

12673-92-6

(coating with, of graphite and titanium electrodes, for brine electrolysis)

TT 7440-32-6, uses and miscellaneous 7782-42-5, uses and miscellaneous

(electrodes, coating of, for brine electrolysis)

L57 ANSWER 6 OF 8 HCA COPYRIGHT 2003 ACS on STN

78:143211 Flexible battery cathode. Hovspian,
Boghos Karnig (du Pont de Nemours, E. I., and Co.). Ger. Offen. DE
2243207 19730308, 20 pp. (German). CODEN: GWXXBX.
APPLICATION: DE 1972-2243207 19720901.

Flexible cathodes for primary cells AB were made by coating slurried sulfides of Fe, Pb, Cd, Cu, or Ni, or oxides of Cu or Fe on 127-203 .mu. thick Pb or Al foils and cold-pressing at 1410-2810 kg/cm2. Thus, FeS prepd. by sintering 1:1 Fe-S mixts. at 450.degree. was sieved, slurried in Cl2FCCF2Cl, spread on a 127 .mu. thick Al foil, and pressed 2 min at 2810 kg/cm2 to give a well adhering coating. A cell made from the coated cathode sheet discharged within 5 min at 125 mA and 1.0 V. 1317-37-9 IT(coating with, on aluminum or lead foil cathodes) 1317-37-9 HCA RNIron sulfide (FeS) (8CI, 9CI) (CA INDEX NAME) CNFe = SIT1317-40-4 (coating with, on lead foil cathodes) 1317-40-4 HCA RNCopper sulfide (CuS) (8CI, 9CI) (CA INDEX NAME) CNCu = S16812-54-7 IT (coating with, on lead foil cathodes) 16812-54-7 HCA RNNickel sulfide (NiS) (8CI, 9CI) (CA INDEX NAME) CNNi = STCH01M 77-2 (Electrochemistry) CC lead cathode battery; aluminum cathode ; iron sulfide coating cathode; cadmium sulfide coating cathode; copper sulfide coating cathode; nickel sulfide coating cathode; oxide copper coating cathode Cathodes IT (battery, aluminum or lead, oxide- or sulfide-coated foils) Coating process IT(of aluminum or lead foil cathodes, with oxides or sulfides) 7429-90-5, uses and miscellaneous 7439-92-1, uses and IT miscellaneous (cathodes, coating of, with sulfides) IT 1317-37-9 (coating with, on aluminum or lead foil cathodes) 1306-23-6, uses and miscellaneous 1309-37-1, uses and ITmiscellaneous 1314-87-0 1317-38-0, uses and miscellaneous

Maples 09/432,334

1317-39-1, uses and miscellaneous 1317-40-4 (coating with, on lead foil cathodes)

IT 16812-54-7

(coating with, on lead foil cathodes)

L57 ANSWER 7 OF 8 HCA COPYRIGHT 2003 ACS on STN 78:105397 Coating of metal halides and oxides with metal sulfides for cathodes. Braun, Leon (Molecular Energy Corp.). Ger. Offen. DE 2135583 19730201, 8 pp. (German). CODEN: GWXXBX. APPLICATION: DE 1971-2135583 19710716.

Cu, Ni, Ag, and Pb halide or oxide particles were coated with Cu, Ni, Ag, and Pb sulfides, resp., by pptn. from a soln. or by spraying. The coated particles were used for battery cathodes low in content of material producing no energy. Thus, 50 g Na2S in 1500 ml EtOH was added to a suspension of 500 g Cu fluoride particles (50-100 mesh) in 1000 ml MeOH to give a Cu sulfide coating. The coated particles were mixed with 3% poly(vinyl chloride) and compacted at .apprx.149.degree. to give a self-supporting elec. conducting cathode of high c.d.

IT 11113-75-0

(coating with, on nickel fluoride, for cathodes)

RN 11113-75-0 HCA

CN Nickel sulfide (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
=======================================	+====================================	+======================================
S	x	7704-34-9
Ni	x	7440-02-0

IT 12751-47-2

(coating with, on silver chloride, for cathodes)

RN 12751-47-2 HCA

CN Silver sulfide (9CI) (CA INDEX NAME)

Component	Ratio	Component Registry Number
===============	+======================================	+======================================
S	x	7704-34-9
Ag	x	7440-22-4

TC H01M

CC 77-2 (Electrochemistry)

Section cross-reference(s): 71

coating metal halide sulfide; copper fluoride sulfide coating; nickel fluoride sulfide coating; silver chloride sulfide coating; lead oxide sulfide coating; cathode battery metal sulfide

IT Cathodes

(battery, metal halides and oxides, coated with sulfides)

IT Coating process

S

```
(of metal halides and oxides, with sulfides, for battery
       cathodes)
    11113-59-0
IT
        (coating of, with copper sulfide, for cathodes)
IT
    1309-60-0
        (coating of, with lead sulfide, for cathodes)
IT
        (coating of, with nickel sulfide, for cathodes)
    7783-90-6
IT
        (coating of, with silver sulfide, for cathodes)
IT
    11115-78-9
        (coating with, on copper fluoride, for cathodes)
    1314-87-0
IT
        (coating with, on lead dioxide, for cathodes)
    11113-75-0
IT
        (coating with, on nickel fluoride, for cathodes)
IT
    12751-47-2
        (coating with, on silver chloride, for cathodes)
    ANSWER 8 OF 8 HCA COPYRIGHT 2003 ACS on STN
76:135112 Electrolysis of brine using coated carbon anodes.
     Chisholm, Raymond S. (PPG Industries, Inc.). U.S. US 3649485
     19720314, 4 pp. (English). CODEN: USXXAM.
                                                 APPLICATION: US
     1968-764618 19681002.
     Coatings of sulfides, nitrides, borides and carbides of Al, Ta, Ti,
AB
     Bi, W, Zr, and Hf mixed with the metals, oxides, sulfides, nitrides,
     borides, and carbides of Au, Ag, Pt, Pd, Ru, Rh, Ir, Os, Ni, Cr, Pb,
     Cu, and Mn are described. For example, a soln. of 3.75 g Pt
     resinate (7.5 wt. Pt), 1 g Ti sulfide and 4 g toluene was painted
     onto a Ti strip which was pickled in HCl soln. prior to painting.
     The painted surface was heated in air to 450.degree. for 1 hr. This
     process was repeated 5 times to yield a tightly bonded coating of Ti
     sulfide-Pt on the Ti base. C electrodes coated in this
     manner are used in cells for prodn. of alkali metal-Cl (both
     diaphragm and Hg-type) and alkali metal-Na chlorate by electrolysis
     of alkali metal chlorides.
     12038-21-0 12673-92-6
IT
        (coating with, of carbon and titanium anodes, for brine
        electrolysis)
     12038-21-0 HCA
RN
     Platinum sulfide (PtS2) (6CI, 7CI, 8CI, 9CI) (CA INDEX NAME)
CN
s = Pt = S
     12673-92-6 HCA
RN
                            (CA INDEX NAME)
     Titanium sulfide (9CI)
CN
                                         Component
                      Ratio
  Component
                                     Registry Number
```

х

7704-34-9

electrolysis)

7440-32-6 Τi C01B; B01K IC 204095000 NCLCC 77 (Electrochemistry) Section cross-reference(s): 49 electrolysis brine coated carbon anode; electrode STcoating electrolysis brine; chlorine prodn elctrolytic cell; alkali metal prodn electrolytic cell; chlorate prodn electrolytic cell IT Brines (electrolysis of, coated carbon anodes for) ITAnodes (for brine electrolysis, coated carbon) Coating process ΙT (of carbon and titanium anodes, for brine electrolysis) 7440-32-6, uses and miscellaneous 7440-44-0, uses and IT miscellaneous (anodes, coated, for brine electrolysis) 7440-06-4, uses and miscellaneous IT (coating with, of carbon and titanium anodes, for brine electrolysis) 11113-84-1 **12038-21-0** 12070-08-5 12673-91-5 IT12673-92-6 (coating with, of carbon and titanium anodes, for brine